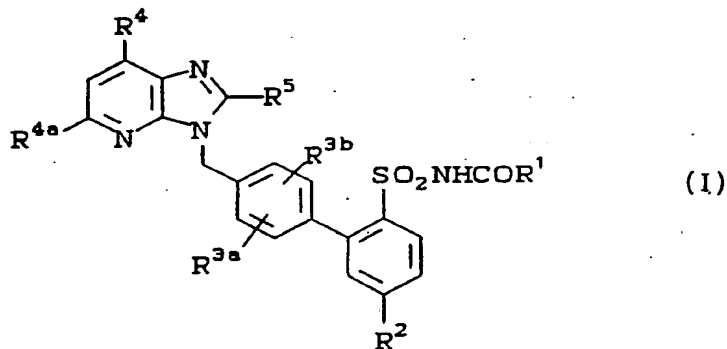




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(54) Title: SUBSTITUTED BIPHENYLMETHYLIMIDAZOPYRIDINES



(57) Abstract

Biphenylmethylimidazopyridines of general structure (I) are angiotensin (II) antagonists and therefore useful in the treatment of hypertension and related cardiovascular disorders and ocular hypertension.

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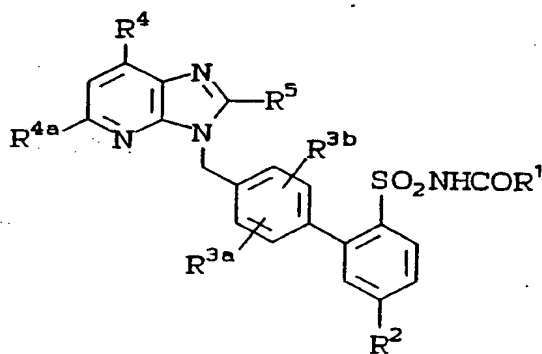
TITLE OF THE INVENTION

SUBSTITUTED BIPHENYLMETHYLMIDAZOPYRIDINES

SUMMARY OF THE INVENTION

This invention is concerned with novel
15 compounds of general structure I:

20



25

I

30

- 2 -

wherein R² is a non-functional substituent such as alkyl, alkoxy, or aryl which are angiotensin II (AII) antagonists demonstrating balanced AT₁/AT₂ activity thus useful in the treatment of hypertension and related cardiovascular disorders and in ocular hypertension.

This invention is also concerned with novel pharmaceutical formulations with one of the novel compounds as active ingredients and the method of treating hypertension and related cardiovascular disorders or ocular hypertension with a novel compound or pharmaceutical formulation thereof.

The invention is also concerned with novel processes for preparing the novel compounds.

BACKGROUND OF THE INVENTION

The renin-angiotensin system (RAS) plays a central role in the regulation of normal blood pressure and seems to be critically involved in hypertension development and maintenance as well as congestive heart failure. Angiotensin II (A II) is an octapeptide hormone produced mainly in the blood during the cleavage of angiotensin I by angiotensin converting enzyme (ACE) localized on the endothelium of blood vessels of lung, kidney, and many other organs. It is the end product of the reninangiotensin system (RAS) and is a powerful arterial vasoconstrictor that exerts its action by interacting with specific receptors present on cell membranes. One of the possible modes of controlling the RAS is angiotensin II receptor antagonism. Several peptide analogs of A II are known to inhibit the effect of

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this hormone by competitively blocking the receptors, but their experimental and clinical applications have been limited by the partial agonist activity and lack of oral absorption [M. Antonaccio. Clin. Exp.

5 Hypertens. A4, 27-46 (1982); D. H. P. Streeten and G. H. Anderson, Jr. - Handbook of Hypertension, Clinical Pharmacology of Antihypertensive Drugs, ed. A. E. Doyle, Vol. 5, pp. 246-271, Elsevier Science Publisher, Amsterdam, The Netherlands, 1984].

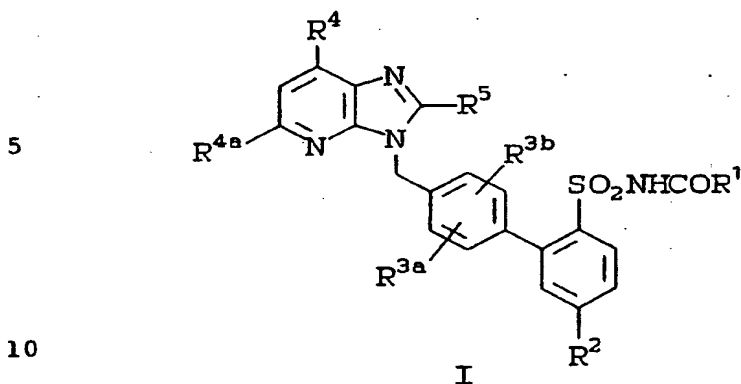
10 Recently, several non-peptide compounds have been described as A II antagonists. Illustrative of such compounds are those disclosed in U.S. Patents 4,207,324; 4,340,598; 4,576,958; 4,582,847; and 4,880,804; in European Patent Applications 028,834;
15 245,637; 253,310; and 291,969; and in articles by A.T. Chiu, et al. [Eur. J. Pharm. Exp. Therap. 157, 13-21 (1988)] and by P.C. Wong, et al. [J. Pharm. Exp. Therap. 247, 1-7(1988)]. All of the U.S. Patents, European Patent Applications 028,834 and
20 253,310 and the two articles disclose substituted imidazole compounds which are generally bonded through a lower alkyl bridge to a substituted phenyl. European Patent Application 399,731 and 400974 disclose imidazopyridines similar to those
25 described herein which are also A-II antagonist.

DETAILED DESCRIPTION OF THE INVENTION

The novel compounds of this invention have structural formula I:

30

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or a pharmaceutically acceptable salt thereof;

15 wherein:

- R¹ is
- a) C₁₋₆ alkyl,
 - b) C₁₋₆ alkylamino,
 - c) C₁₋₆ alkoxy-(CH₂)_n-, wherein n is 1 or 2,
 - d) aryl-(CH₂)_s-, wherein s is 0 to 3
 - e) C₁₋₆ alkylthio-(CH₂)_n-,
 - f) aryl, either unsubstituted or substituted with
 - 1) C₁₋₆ alkyl,
 - 2) aryloxy,
 - 3) C₁₋₆ alkoxy,
 - 4) -Cl,
 - 5) -Br, or
 - 6) C₁₋₆ alkylamino;
- 20
- 25
- 30

- R² is
- a) -Cl,
 - b) C₁₋₆ alkyl,

- 5 -

- c) C₁₋₅ alkoxy,
d) C₁₋₅ alkoxy-CH₂-,
e) di(C₁₋₅ alkyl)amino-CH₂-,
f) pyrrolidin-1-yl-CH₂-,
g) morpholin-1-yl-CH₂-,
h) polyfluoro-C₁₋₅ alkoxy,
i) aryl,
j) C₁₋₅ alkyl-S-(O)_S-(CH₂)_S-or
k) aryl-(CH₂)_n-;

R^{3a} and R^{3b} are independently

- a) H,
b) F, Cl, Br or I,
c) C₁₋₄ alkyl,
d) C₁₋₄ alkoxy, or
e) aryl;

R^{3a} and R^{3b} on adjacent carbons can be joined together to form a benzo group;

R⁴ and R^{4a} are independently

- a) C₁₋₃ alkyl,
b) polyfluoro-C₁₋₃ alkyl,
c) -CONHR¹,
d) -CO₂R¹ or
e) -CONH (CH₂)_n-aryl;

not H

R⁵ is hydrogen or C₁₋₅ alkyl;

In the above definitions, aryl is meant to include phenyl, naphthyl and 2-, 3-, or 4-pyridyl.

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The terms "alkyl" and "alkoxy", include both straight - and branched-chain groups where the number of carbons permit.

5 One embodiment of the novel compounds is that wherein R^4 and R^{4a} are both C_{1-3} alkyl, especially methyl, and R^5 is C_{1-5} alkyl, especially ethyl.

10 A class of compounds within this embodiment is that wherein R^2 is C_{1-6} alkyl, especially n-propyl.

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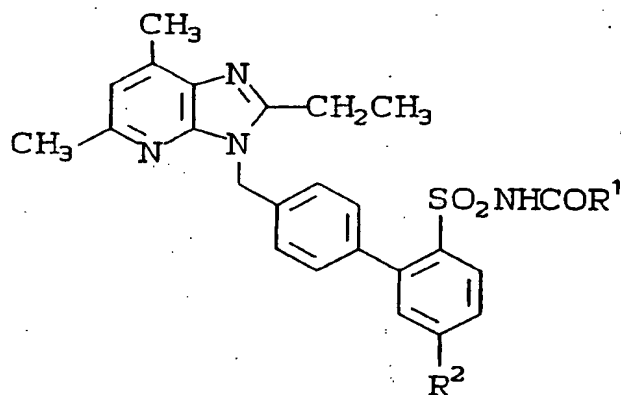
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Specific compounds exemplifying the novel compounds of this invention are described in Table I.

TABLE I



	#(EX)	SO ₂ NHCOR ¹	R ²
20	1	SO ₂ NHCO(CH ₂) ₄ CH ₃	CH ₃
	2	SO ₂ NHCO(CH ₂) ₄ CH ₃	CH ₂ CH ₃
	3 (1)	SO ₂ NHCO(CH ₂) ₄ CH ₃	(CH ₂) ₂ CH ₃
	4	SO ₂ NHCO(CH ₂) ₄ CH ₃	(CH ₂) ₃ CH ₃
	5 (8)	SO ₂ NHCO(CH ₂) ₄ CH ₃	CH(CH ₃) ₂
25	6 (13)	SO ₂ NHCO(CH ₂) ₄ CH ₃	O(CH ₂) ₃ CH ₃
	7	SO ₂ NHCO(CH ₂) ₄ CH ₃	CH ₂ CH(CH ₃) ₂
	8	SO ₂ NHCO(CH ₂) ₄ CH ₃	OCH ₃
	9	SO ₂ NHCO(CH ₂) ₄ CH ₃	CH ₂ SCH ₃
	10	SO ₂ NHCO(CH ₂) ₄ CH ₃	CH ₂ OCH ₃
30	11	SO ₂ NHCO(CH ₂) ₄ CH ₃	OCH ₂ CH ₃
	12	SO ₂ NHCO(CH ₂) ₄ CH ₃	Ph

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#(EX)	$\text{SO}_2\text{NHCOR}^1$	R^2
5	13 $\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{C}(\text{CH}_3)_2\text{CH}_2\text{CH}_3$
	14 $\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{C}(\text{CH}_3)_3$
	15 (6) $\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_3)_2$
	16 (11) $\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2$
	17 (12) $\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2\text{O}$
10	18 $\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	OCH_2CF_3
	19 $\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{OCH}(\text{CH}_3)_2$
	20 $\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	SCH_2CH_3
	21 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	CH_3
	22 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	CH_2CH_3
15	23 (10) $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$(\text{CH}_2)_2\text{CH}_3$
	24 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$(\text{CH}_2)_3\text{CH}_3$
	25 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}(\text{CH}_3)_2$
	26 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{O}(\text{CH}_2)_3\text{CH}_3$
	27 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{CH}(\text{CH}_3)_2$
20	28 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	OCH_3
	29 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	CH_2SCH_3
	30 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	CH_2OCH_3
	31 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	OCH_2CH_3
	32 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	Ph
25	33 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{C}(\text{CH}_3)_2\text{CH}_2\text{CH}_3$
	34 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{C}(\text{CH}_3)_3$
	35 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_3)_2$
	36 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2$
	37 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2\text{O}$
30	38 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	OCH_2CF_3
	39 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{OCH}(\text{CH}_3)_2$
	40 $\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	SCH_2CH_3
	41 $\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	CH_3
	42 $\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	CH_2CH_3
	43 (9) $\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	$(\text{CH}_2)_2\text{CH}_3$

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	#(EX)	SO ₂ NHCOR ¹	R ²
	44	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	(CH ₂) ₃ CH ₃
	45	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	CH(CH ₃) ₂
5	46	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	O(CH ₂) ₃ CH ₃
	47	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	CH ₂ CH(CH ₃) ₂
	48	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	OCH ₃
	49	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	CH ₂ SCH ₃
	50	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	CH ₂ OCH ₃
10	51	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	OCH ₂ CH ₃
	52	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	Ph
	53	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	C(CH ₃) ₂ CH ₂ CH ₃
	54	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	C(CH ₃) ₃
	55	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	CH ₂ N(CH ₃) ₂
15	56	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	CH ₂ N(CH ₂ CH ₂) ₂
	57	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	CH ₂ N(CH ₂ CH ₂) ₂ O
	58	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	OCH ₂ CF ₃
	59	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	OCH(CH ₃) ₂
	60	SO ₂ NHCONH(CH ₂) ₃ CH ₃	SCH ₂ CH ₃
20	61	SO ₂ NHCONH(CH ₂) ₃ CH ₃	CH ₃
	62	SO ₂ NHCONH(CH ₂) ₃ CH ₃	CH ₂ CH ₃
	63 (2)	SO ₂ NHCONH(CH ₂) ₃ CH ₃	(CH ₂) ₂ CH ₃
	64	SO ₂ NHCONH(CH ₂) ₃ CH ₃	(CH ₂) ₃ CH ₃
	65	SO ₂ NHCONH(CH ₂) ₃ CH ₃	CH(CH ₃) ₂
25	66 (7)	SO ₂ NHCONH(CH ₂) ₃ CH ₃	O(CH ₂) ₃ CH ₃
	67	SO ₂ NHCONH(CH ₂) ₃ CH ₃	CH ₂ CH(CH ₃) ₂
	68	SO ₂ NHCONH(CH ₂) ₃ CH ₃	OCH ₃
	69	SO ₂ NHCONH(CH ₂) ₃ CH ₃	CH ₂ SCH ₃
	70	SO ₂ NHCONH(CH ₂) ₃ CH ₃	CH ₂ OCH ₃
30	71	SO ₂ NHCONH(CH ₂) ₃ CH ₃	OCH ₂ CH ₃
	72	SO ₂ NHCONH(CH ₂) ₃ CH ₃	Ph
	73	SO ₂ NHCONH(CH ₂) ₃ CH ₃	C(CH ₃) ₂ CH ₂ CH ₃
	74	SO ₂ NHCONH(CH ₂) ₃ CH ₃	C(CH ₃) ₃

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#(EX)	$\text{SO}_2\text{NHCOR}^1$	R^2
5	75 $\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_3)_2$
	76 $\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2$
	77 $\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2\text{O}$
	78 $\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	OCH_2CF_3
	79 $\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	$\text{OCH}(\text{CH}_3)_2$
10	80 $\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	SCH_2CH_3
	81 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	CH_3
	82 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	CH_2CH_3
	83 (3) $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$(\text{CH}_2)_2\text{CH}_3$
	84 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$(\text{CH}_2)_3\text{CH}_3$
15	85 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{CH}(\text{CH}_3)_2$
	86 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{O}(\text{CH}_2)_3\text{CH}_3$
	87 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{CH}_2\text{CH}(\text{CH}_3)_2$
	88 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	OCH_3
	89 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	CH_2SCH_3
20	90 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	CH_2OCH_3
	91 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	OCH_2CH_3
	92 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	Ph
	93 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{C}(\text{CH}_3)_2\text{CH}_2\text{CH}_3$
	94 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{C}(\text{CH}_3)_3$
25	95 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{CH}_2\text{N}(\text{CH}_3)_2$
	96 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2$
	97 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2\text{O}$
	98 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	OCH_2CF_3
	99 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{OCH}(\text{CH}_3)_2$
30	100 $\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	SCH_2CH_3
	101 $\text{SO}_2\text{NHCO}(2\text{-PhO})\text{Ph}$	CH_3
	102 $\text{SO}_2\text{NHCO}(2\text{-PhO})\text{Ph}$	CH_2CH_3
	103 (4) $\text{SO}_2\text{NHCO}(2\text{-PhO})\text{Ph}$	$(\text{CH}_2)_2\text{CH}_3$
	104 $\text{SO}_2\text{NHCO}(2\text{-PhO})\text{Ph}$	$(\text{CH}_2)_2\text{CH}_3$
	105 $\text{SO}_2\text{NHCO}(2\text{-PhO})\text{Ph}$	$\text{CH}(\text{CH}_3)_2$

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	#(EX)	SO ₂ NHCOR ¹	R ²
	106	SO ₂ NHCO(2-PhO)Ph	O(CH ₂) ₃ CH ₃
	107	SO ₂ NHCO(2-PhO)Ph	CH ₂ CH(CH ₃) ₂
5	108	SO ₂ NHCO(2-PhO)Ph	OCH ₃
	109	SO ₂ NHCO(2-PhO)Ph	CH ₂ SCH ₃
	110	SO ₂ NHCO(2-PhO)Ph	CH ₂ OCH ₃
	111	SO ₂ NHCO(2-PhO)Ph	OCH ₂ CH ₃
	112	SO ₂ NHCO(2-PhO)Ph	Ph
10	113	SO ₂ NHCO(2-PhO)Ph	C(CH ₃) ₂ CH ₂ CH ₃
	114	SO ₂ NHCO(2-PhO)Ph	C(CH ₃) ₃
	115	SO ₂ NHCO(2-PhO)Ph	CH ₂ N(CH ₃) ₂
	116	SO ₂ NHCO(2-PhO)Ph	CH ₂ N(CH ₂ CH ₂) ₂
	117	SO ₂ NHCO(2-PhO)Ph	CH ₂ N(CH ₂ CH ₂) ₂ O
15	118	SO ₂ NHCO(2-PhO)Ph	OCH ₂ CF ₃
	119	SO ₂ NHCO(2-PhO)Ph	OCH(CH ₃) ₂
	120	SO ₂ NHCO(2-PhO)Ph	SCH ₂ CH ₃
	121	SO ₂ NHCO(2-EtO)Ph	CH ₃
	122	SO ₂ NHCO(2-EtO)Ph	CH ₂ CH ₃
20	123	SO ₂ NHCO(2-EtO)Ph	(CH ₂) ₂ CH ₃
	124	SO ₂ NHCO(2-EtO)Ph	(CH ₂) ₂ CH ₃
	125	SO ₂ NHCO(2-EtO)Ph	CH(CH ₃) ₂
	126	SO ₂ NHCO(2-EtO)Ph	O(CH ₂) ₃ CH ₃
	127	SO ₂ NHCO(2-EtO)Ph	CH ₂ CH(CH ₃) ₂
25	128	SO ₂ NHCO(2-EtO)Ph	OCH ₃
	129	SO ₂ NHCO(2-EtO)Ph	CH ₂ SCH ₃
	130	SO ₂ NHCO(2-EtO)Ph	CH ₂ OCH ₃
	131	SO ₂ NHCO(2-EtO)Ph	OCH ₂ CH ₃
	132	SO ₂ NHCO(2-EtO)Ph	Ph
30	133	SO ₂ NHCO(2-EtO)Ph	C(CH ₃) ₂ CH ₂ CH ₃
	134	SO ₂ NHCO(2-EtO)Ph	C(CH ₃) ₃
	135	SO ₂ NHCO(2-EtO)Ph	CH ₂ N(CH ₃) ₂
	136	SO ₂ NHCO(2-EtO)Ph	CH ₂ N(CH ₂ CH ₂) ₂

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	#(EX)	SO ₂ NHCOR ¹	R ²
	137	SO ₂ NHCO(2-EtO)Ph	CH ₂ N(CH ₂ CH ₂) ₂ O
	138	SO ₂ NHCO(2-EtO)Ph	OCH ₂ CF ₃
5	139	SO ₂ NHCO(2-EtO)Ph	OCH(CH ₃) ₂
	140	SO ₂ NHCO(2-EtO)Ph	SCH ₂ CH ₃
	141	SO ₂ NHCOPh	CH ₃
	142	SO ₂ NHCOPh	CH ₂ CH ₃
	143 (5)	SO ₂ NHCOPh	(CH ₂) ₂ CH ₃
10	144	SO ₂ NHCOPh	(CH ₂) ₂ CH ₃
	145	SO ₂ NHCOPh	CH(CH ₃) ₂
	146	SO ₂ NHCOPh	O(CH ₂) ₃ CH ₃
	147	SO ₂ NHCOPh	CH ₂ N(CH ₃) ₂
	148	SO ₂ NHCOPh	OCH ₃
15	149	SO ₂ NHCOPh	CH ₂ SCH ₃
	150	SO ₂ NHCOPh	CH ₂ OCH ₃
	151	SO ₂ NHCOPh	OCH ₂ CH ₃
	152	SO ₂ NHCOPh	Ph
	153	SO ₂ NHCOPh	C(CH ₃) ₂ CH ₂ CH ₃
20	154	SO ₂ NHCOPh	C(CH ₃) ₃
	155	SO ₂ NHCOPh	CH ₂ N(CH ₃) ₂
	156	SO ₂ NHCOPh	CH ₂ N(CH ₂ CH ₂) ₂
	157	SO ₂ NHCOPh	CH ₂ N(CH ₂ CH ₂) ₂ O
	158	SO ₂ NHCOPh	OCH ₂ CF ₃
25	159	SO ₂ NHCOPh	OCH(CH ₃) ₂
	160	SO ₂ NHCOPh	SCH ₂ CH ₃
	161	SO ₂ NHCO(CH ₂) ₄ CH ₃	CH ₃
	162	SO ₂ NHCO(CH ₂) ₄ CH ₃	CH ₂ CH ₃
	163	SO ₂ NHCO(CH ₂) ₄ CH ₃	(CH ₂) ₂ CH ₃
30	164	SO ₂ NHCO(CH ₂) ₄ CH ₃	(CH ₂) ₂ CH ₃
	165	SO ₂ NHCO(CH ₂) ₄ CH ₃	CH(CH ₃) ₂
	166	SO ₂ NHCO(CH ₂) ₄ CH ₃	O(CH ₂) ₃ CH ₃
	167	SO ₂ NHCO(CH ₂) ₄ CH ₃	CH ₂ N(CH ₃) ₂

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	#(EX)	SO ₂ NHCO ¹	R ²
	168	SO ₂ NHCO(CH ₂) ₄ CH ₃	OCH ₃
	169	SO ₂ NHCO(CH ₂) ₄ CH ₃	CH ₂ SCH ₃
5	170	SO ₂ NHCO(CH ₂) ₄ CH ₃	CH ₂ OCH ₃
	171	SO ₂ NHCO(CH ₂) ₄ CH ₃	OCH ₂ CH ₃
	172	SO ₂ NHCO(CH ₂) ₄ CH ₃	Ph
	173	SO ₂ NHCO(CH ₂) ₄ CH ₃	C(CH ₃) ₂ CH ₂ CH ₃
	174	SO ₂ NHCO(CH ₂) ₄ CH ₃	C(CH ₃) ₃
10	175	SO ₂ NHCO(CH ₂) ₄ CH ₃	CH ₂ N(CH ₃) ₂
	176	SO ₂ NHCO(CH ₂) ₄ CH ₃	CH ₂ N(CH ₂ CH ₂) ₂
	177	SO ₂ NHCO(CH ₂) ₄ CH ₃	CH ₂ N(CH ₂ CH ₂) ₂ O
	178	SO ₂ NHCO(CH ₂) ₄ CH ₃	OCH ₂ CF ₃
	179	SO ₂ NHCO(CH ₂) ₄ CH ₃	OCH(CH ₃) ₂
15	180	SO ₂ NHCO(CH ₂) ₄ CH ₃	SCH ₂ CH ₃
	181	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	CH ₃
	182	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	CH ₂ CH ₃
	183	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	(CH ₂) ₂ CH ₃
	184	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	(CH ₂) ₂ CH ₃
20	185	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	CH(CH ₃) ₂
	186	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	O(CH ₂) ₃ CH ₃
	187	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	CH ₂ N(CH ₃) ₂
	188	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	OCH ₃
	189	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	CH ₂ SCH ₃
25	190	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	CH ₂ OCH ₃
	191	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	OCH ₂ CH ₃
	192	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	Ph
	193	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	C(CH ₃) ₂ CH ₂ CH ₃
	194	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	C(CH ₃) ₃
30	195	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	CH ₂ N(CH ₃) ₂
	196	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	CH ₂ N(CH ₂ CH ₂) ₂
	197	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	CH ₂ N(CH ₂ CH ₂) ₂ O
	198	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	OCH ₂ CF ₃

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	#(EX)	SO ₂ NHCOR ¹	R ²
	199	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	OCH(CH ₃) ₂
	200	SO ₂ NHCOCH ₂ O(CH ₂) ₂ CH ₃	SCH ₂ CH ₃
5	201	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	CH ₃
	202	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	CH ₂ CH ₃
	203	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	(CH ₂) ₂ CH ₃
	204	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	(CH ₂) ₃ CH ₃
	205	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	CH(CH ₃) ₂
10	206	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	O(CH ₂) ₃ CH ₃
	207	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	CH ₂ CH(CH ₃) ₂
	208	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	OCH ₃
	209	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	CH ₂ SCH ₃
	210	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	CH ₂ OCH ₃
15	211	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	OCH ₂ CH ₃
	212	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	Ph
	213	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	C(CH ₃) ₂ CH ₂ CH ₃
	214	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	C(CH ₃) ₃
	215	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	CH ₂ N(CH ₃) ₂
20	216	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	CH ₂ N(CH ₂ CH ₂) ₂
	217	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	CH ₂ N(CH ₂ CH ₂) ₂ O
	218	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	OCH ₂ CF ₃
	219	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	OCH(CH ₃) ₂
	220	SO ₂ NHCOCH ₂ OCH ₂ CH ₃	SCH ₂ CH ₃
25	221	SO ₂ NHCONH(CH ₂) ₃ CH ₃	CH ₃
	222	SO ₂ NHCONH(CH ₂) ₃ CH ₃	CH ₂ CH ₃
	223	SO ₂ NHCONH(CH ₂) ₃ CH ₃	(CH ₂) ₂ CH ₃
	224	SO ₂ NHCONH(CH ₂) ₃ CH ₃	(CH ₂) ₃ CH ₃
	225	SO ₂ NHCONH(CH ₂) ₃ CH ₃	CH(CH ₃) ₂
30	226	SO ₂ NHCONH(CH ₂) ₃ CH ₃	O(CH ₂) ₃ CH ₃
	227	SO ₂ NHCONH(CH ₂) ₃ CH ₃	CH ₂ CH(CH ₃) ₂
	228	SO ₂ NHCONH(CH ₂) ₃ CH ₃	OCH ₃
	229	SO ₂ NHCONH(CH ₂) ₃ CH ₃	CH ₂ SCH ₃
	230	SO ₂ NHCONH(CH ₂) ₃ CH ₃	CH ₂ OCH ₃

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	#(EX)	SO ₂ NHCOR ¹	R ²
	231	SO ₂ NHCONH(CH ₂) ₃ CH ₃	OCH ₂ CH ₃
	232	SO ₂ NHCONH(CH ₂) ₃ CH ₃	Ph
5	233	SO ₂ NHCONH(CH ₂) ₃ CH ₃	C(CH ₃) ₂ CH ₂ CH ₃
	234	SO ₂ NHCONH(CH ₂) ₃ CH ₃	C(CH ₃) ₃
	235	SO ₂ NHCON(CH ₂) ₃ CH ₃	CH ₂ N(CH ₃) ₂
	236	SO ₂ NHCON(CH ₂) ₃ CH ₃	CH ₂ N(CH ₂ CH ₂) ₂
	236	SO ₂ NHCON(CH ₂) ₃ CH ₃	CH ₂ N(CH ₂ CH ₂) ₂ O
10	238	SO ₂ NHCON(CH ₂) ₃ CH ₃	OCH ₂ CF ₃
	239	SO ₂ NHCON(CH ₂) ₃ CH ₃	OCH(CH ₃) ₂
	240	SO ₂ NHCON(CH ₂) ₃ CH ₃	SCH ₂ CH ₃
	241	SO ₂ NHCO(CH ₂) ₂ Ph	CH ₃
	242	SO ₂ NHCO(CH ₂) ₂ Ph	CH ₂ CH ₃
15	243	SO ₂ NHCO(CH ₂) ₂ Ph	(CH ₂) ₂ CH ₃
	244	SO ₂ NHCO(CH ₂) ₂ Ph	(CH ₂) ₃ CH ₃
	245	SO ₂ NHCO(CH ₂) ₂ Ph	CH(CH ₃) ₂
	246	SO ₂ NHCO(CH ₂) ₂ Ph	O(CH ₂) ₃ CH ₃
	247	SO ₂ NHCO(CH ₂) ₂ Ph	CH ₂ CH(CH ₃) ₂
20	248	SO ₂ NHCO(CH ₂) ₂ Ph	OCH ₃
	249	SO ₂ NHCO(CH ₂) ₂ Ph	CH ₂ SCH ₃
	250	SO ₂ NHCO(CH ₂) ₂ Ph	CH ₂ OCH ₃
	251	SO ₂ NHCO(CH ₂) ₂ Ph	OCH ₂ CH ₃
	252	SO ₂ NHCO(CH ₂) ₂ Ph	Ph
25	253	SO ₂ NHCO(CH ₂) ₂ Ph	C(CH ₃) ₂ CH ₂ CH ₃
	254	SO ₂ NHCO(CH ₂) ₂ Ph	C(CH ₃) ₃
	255	SO ₂ NHCO(CH ₂) ₂ Ph	CH ₂ N(CH ₃) ₂
	256	SO ₂ NHCO(CH ₂) ₂ Ph	CH ₂ N(CH ₂ CH ₂) ₂
	257	SO ₂ NHCO(CH ₂) ₂ Ph	CH ₂ N(CH ₂ CH ₂) ₂ O
30	258	SO ₂ NHCO(CH ₂) ₂ Ph	OCH ₂ CF ₃
	259	SO ₂ NHCO(CH ₂) ₂ Ph	OCH(CH ₃) ₂
	260	SO ₂ NHCO(CH ₂) ₂ Ph	SCH ₂ CH ₃

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#(EX)	$\text{SO}_2\text{NHCOR}^1$	R^2
5	261 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	CH_3
	262 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	CH_2CH_3
	263 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	$(\text{CH}_2)_2\text{CH}_3$
	264 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	$(\text{CH}_2)_2\text{CH}_3$
	265 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	$\text{CH}(\text{CH}_3)_2$
10	266 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	$\text{O}(\text{CH}_2)_3\text{CH}_3$
	267 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	$\text{CH}_2\text{CH}(\text{CH}_3)_2$
	268 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	OCH_3
	269 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	CH_2SCH_3
	270 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	CH_2OCH_3
15	271 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	OCH_2CH_3
	272 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	Ph
	273 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	$\text{C}(\text{CH}_3)_2\text{CH}_2\text{CH}_3$
	274 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	$\text{C}(\text{CH}_3)_3$
	275 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	$\text{CH}_2\text{N}(\text{CH}_3)_2$
20	276 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2$
	277 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2\text{O}$
	278 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	OCH_2CF_3
	279 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	$\text{OCH}(\text{CH}_3)_2$
	280 $\text{SO}_2\text{NHCOR}(2\text{-PhO})\text{Ph}$	SCH_2CH_3
25	281 $\text{SO}_2\text{NHCOR}(2\text{-EtO})\text{Ph}$	CH_3
	282 $\text{SO}_2\text{NHCOR}(2\text{-EtO})\text{Ph}$	CH_2CH_3
	283 $\text{SO}_2\text{NHCOR}(2\text{-EtO})\text{Ph}$	$(\text{CH}_2)_2\text{CH}_3$
	284 $\text{SO}_2\text{NHCOR}(2\text{-EtO})\text{Ph}$	$(\text{CH}_2)_2\text{CH}_3$
	285 $\text{SO}_2\text{NHCOR}(2\text{-EtO})\text{Ph}$	$\text{CH}(\text{CH}_3)_2$
30	286 $\text{SO}_2\text{NHCOR}(2\text{-EtO})\text{Ph}$	$\text{O}(\text{CH}_2)_3\text{CH}_3$
	287 $\text{SO}_2\text{NHCOR}(2\text{-EtO})\text{Ph}$	$\text{CH}_2\text{CH}(\text{CH}_3)_2$
	288 $\text{SO}_2\text{NHCOR}(2\text{-EtO})\text{Ph}$	OCH_3
	289 $\text{SO}_2\text{NHCOR}(2\text{-EtO})\text{Ph}$	CH_2SCH_3
	290 $\text{SO}_2\text{NHCOR}(2\text{-EtO})\text{Ph}$	CH_2OCH_3
	291 $\text{SO}_2\text{NHCOR}(2\text{-EtO})\text{Ph}$	OCH_2CH_3

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	<u>#(EX)</u>	<u>SO₂NHCOR¹</u>	<u>R²</u>
	293	SO ₂ NHCO(2-EtO)Ph	Ph
	294	SO ₂ NHCO(2-EtO)Ph	C(CH ₃) ₂ CH ₂ CH ₃
5	295	SO ₂ NHCO(2-EtO)Ph	C(CH ₃) ₃
	296	SO ₂ NHCO(2-EtO)Ph	CH ₂ N(CH ₃) ₂
	297	SO ₂ NHCO(2-EtO)Ph	CH ₂ N(CH ₂ CH ₂) ₂
	298	SO ₂ NHCO(2-PhO)Ph	OCH ₂ CF ₃
	299	SO ₂ NHCO(2-PhO)Ph	OCH(CH ₃) ₂
10	300	SO ₂ NHCO(2-PhO)Ph	SCH ₂ CH ₃
	301	SO ₂ NHCOPh	CH ₃
	302	SO ₂ NHCOPh	CH ₂ CH ₃
	303	SO ₂ NHCOPh	(CH ₂) ₂ CH ₃
	304	SO ₂ NHCOPh	(CH ₂) ₂ CH ₃
15	305	SO ₂ NHCOPh	CH(CH ₃) ₂
	306	SO ₂ NHCOPh	O(CH ₂) ₃ CH ₃
	307	SO ₂ NHCOPh	CH ₂ N(CH ₃) ₂
	308	SO ₂ NHCOPh	OCH ₃
	309	SO ₂ NHCOPh	CH ₂ SCH ₃
20	310	SO ₂ NHCOPh	CH ₂ OCH ₃
	311	SO ₂ NHCOPh	OCH ₂ CH ₃
	312	SO ₂ NHCOPh	Ph
	313	SO ₂ NHCOPh	C(CH ₃) ₂ CH ₂ CH ₃
	314	SO ₂ NHCOPh	C(CH ₃) ₃
25	315	SO ₂ NHCOPh	CH ₂ N(CH ₃) ₂
	316	SO ₂ NHCOPh	CH ₂ N(CH ₂ CH ₂) ₂
	317	SO ₂ NHCOPh	CH ₂ N(CH ₂ CH ₂) ₂ O
	318	SO ₂ NHCOPh	OCH ₂ CF ₃
	319	SO ₂ NHCOPh	OCH(CH ₃) ₂
30	320	SO ₂ NHCOPh	SCH ₂ CH ₃

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Scheme Descriptions:

The general procedure used to prepare many of the 5'-substituted derivatives is illustrated in Scheme I. Commercially available 4-substituted benzenesulfonyl chlorides ($R^2 = i\text{-Pr}, n\text{-BuO}, \text{tert-amyl}, \text{Me}, \text{Et}, n\text{-Pr}, t\text{-Bu}$) are reacted with $t\text{-BuNH}_2$ in CH_2Cl_2 or CHCl_3 to provide derivative 2 in good yield. Dianion generation in THF, with 2.5 equivalents of $n\text{-BuLi}$, followed by quench with triisopropyl borate provides boric acid derivative 3 in excellent yield, after hydrolysis with dilute acid. Palladium catalyzed coupling of boric acid 3 and 4-bromobenzyl derivative 4 in the presence of 1.25 N NaOH, EtOH and toluene, affords an excellent yield of the desired coupled product. Deprotection using TFA is followed by coupling using methods A or B with the appropriate acid or acid chloride to prepare acylsulfonamides and method C with the appropriate isocyanate to prepare sulfonylureas.

When the desired 4-substituted benzenesulfonyl chlorides are not commercially available, the necessary 4-substituted benzenet-butylsulfonamide (2) derivatives can be prepared using a variety of procedures. These procedures are outlined in Scheme II (A - F). In Scheme II A, a variety of trimethylstannyl derivatives could be coupled to aryl bromide 8 in the presence of $\text{Pd(PPh}_3)_4$ or $\text{PdCl}_2(\text{PPh}_3)_2$. In this example this is followed by hydrogenation to obtain the isobutyl derivative. Introduction of an aryl ring is best accomplished using the palladium catalyzed boric acid coupling method illustrated in Scheme II B. Thiomethyl and aminomethyl derivatives are both

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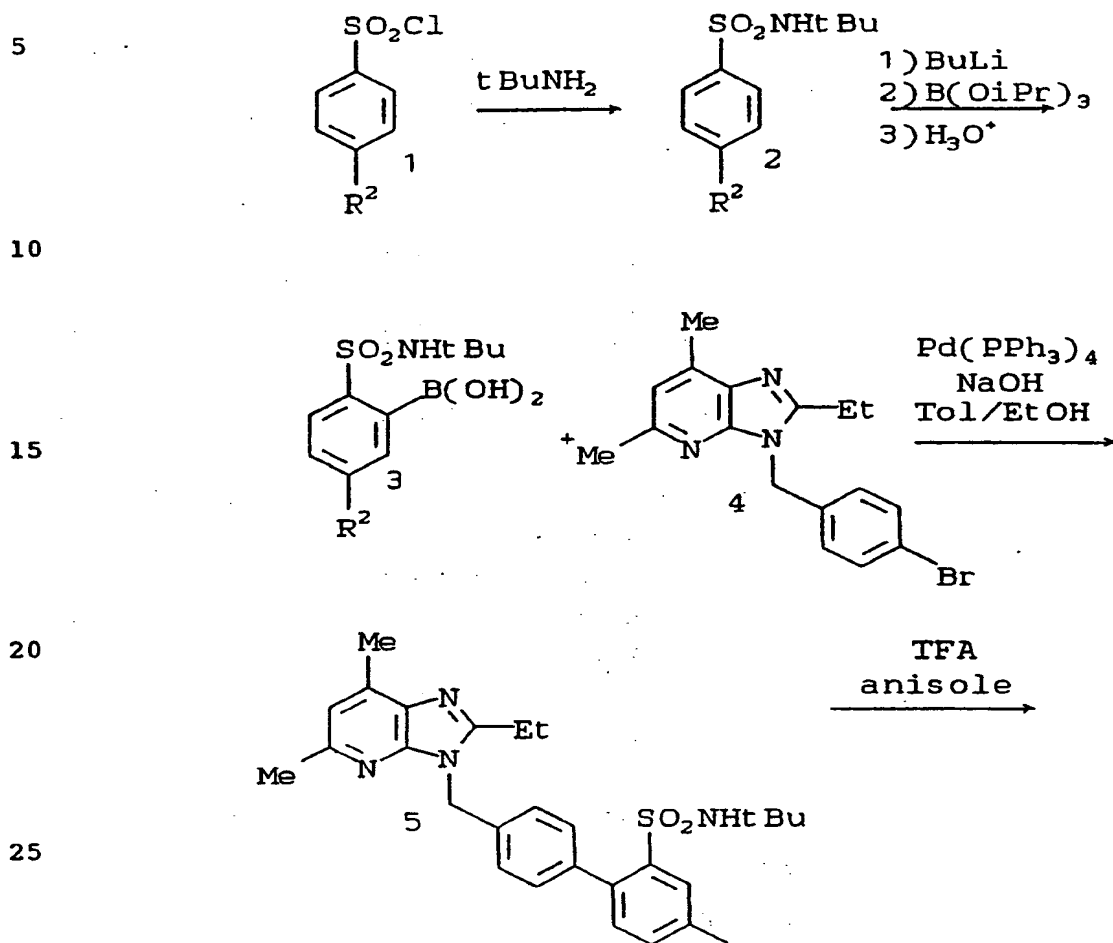
prepared from bromomethyl derivative 11 as illustrated in Schemes II C and D receptively. A convenient manner to introduce alkoxy substituents is illustrated in Scheme II E. An alternative procedure to prepare alkyl derivatives starting with an alkylbenzene is illustrated in Scheme II F.

Antagonists with 5'-alkoxy methyl derivatives are best prepared using the protocol outlined in Scheme III. Palladium catalyzed coupling of 5-methyl-2-t-butylsulfonamide phenylboric acid with methyl 4-iodobenzoate affords derivative 12. Benzylic bromination, utilizing NBS in refluxing CCl_4 , with a catalytic amount of AIBN or alternatively, benzoyl peroxide, provides the desired bromomethyl derivative that is then reacted with the appropriate sodium alkoxide to afford derivative 14. Reduction of the ester to the primary alcohol with LAH is followed by conversion to the bromomethyl derivative (16) with PBr_3 . Alkylation of the sodium salt of the heterocycle with 16 in DMF provides derivative 6 ($\text{R}^2 = \text{CH}_2\text{OR}$). The antagonist is completed as previously illustrated in Scheme I.

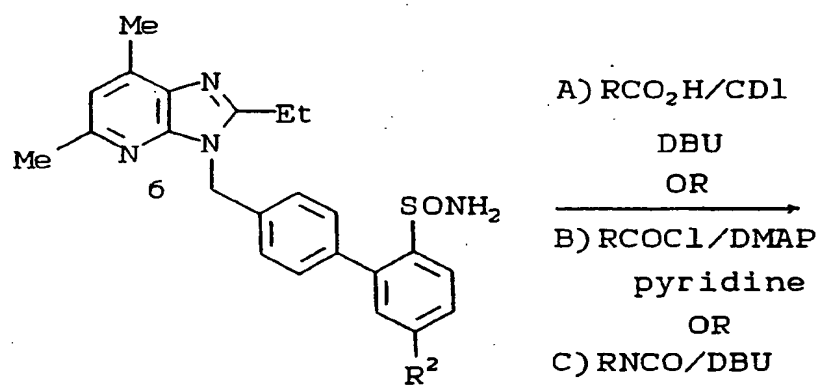
Introduction of substituents into the central phenyl of the biphenyl moiety is best accomplished using the protocol illustrated in scheme IV. Palladium catalyzed coupling of 5-substituted-2-t-butyl-sulfonamide phenylboric acid with a substituted methyl 4-iodobenzoate or bromobenzoate affords derivative 17. Reduction of the ester to the primary alcohol with LAH is followed by conversion to the bromomethyl derivative (19) with PBr_3 . Alkylation of the sodium salt of the heterocycle with 19 in DMF provides derivative 6. The antagonist is completed as previously illustrated in Scheme I.

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SCHEME I

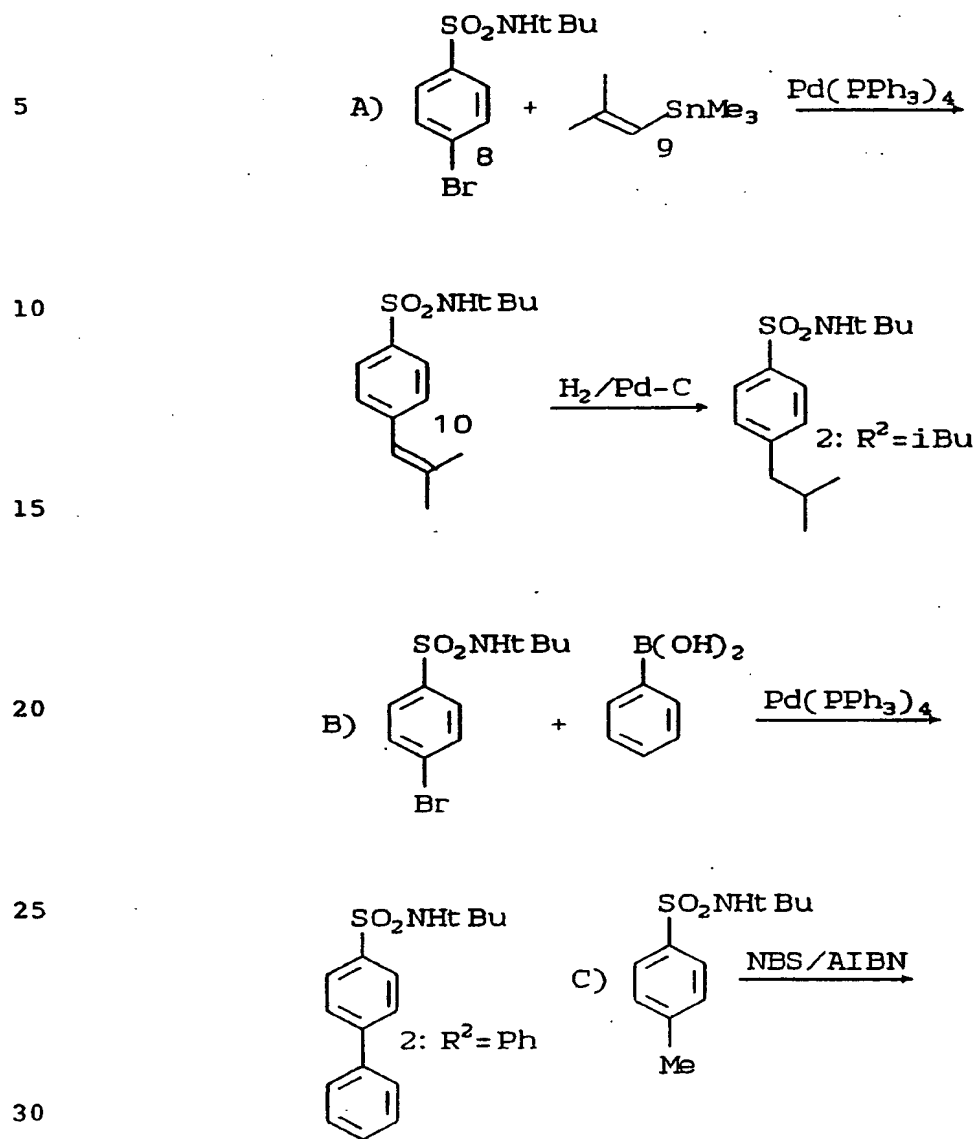


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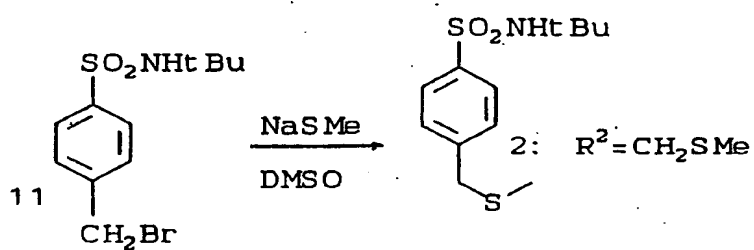
SCHEME I Cont'd.

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SCHEME II



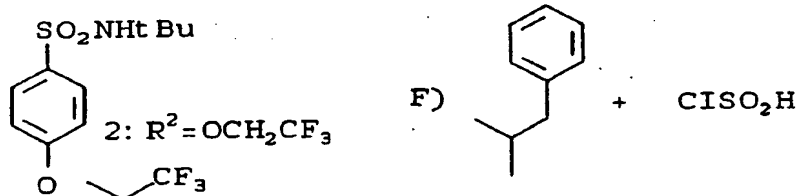
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SCHEME II Cont'd.

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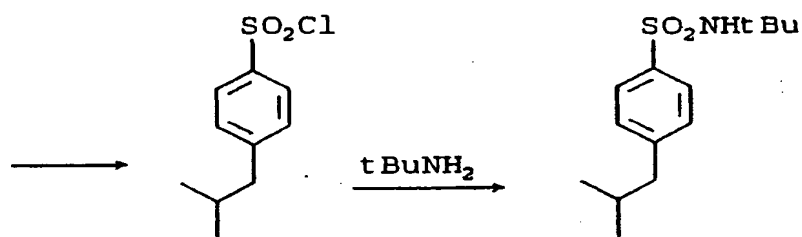
SCHEME II Cont'd.

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SCHEME III

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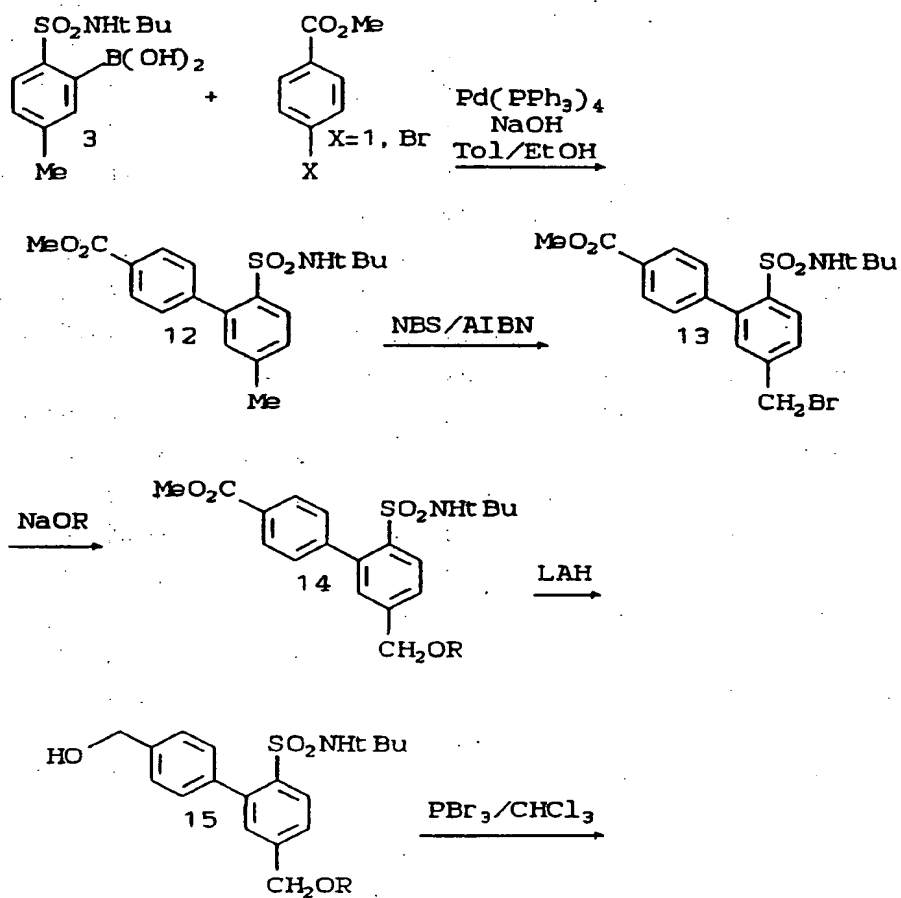
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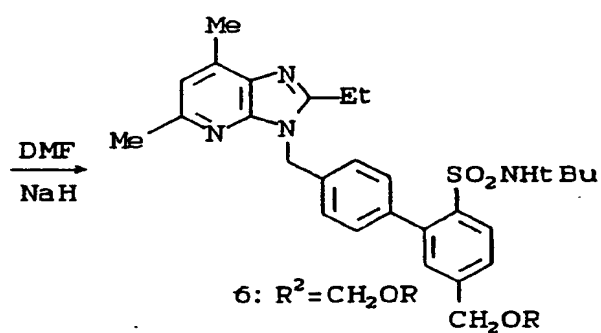
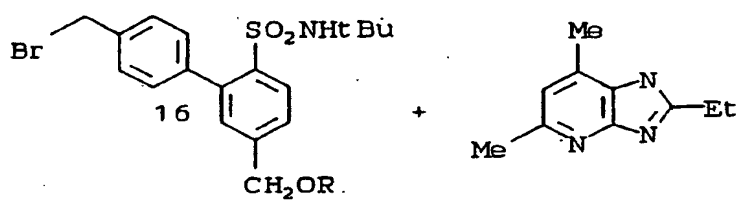
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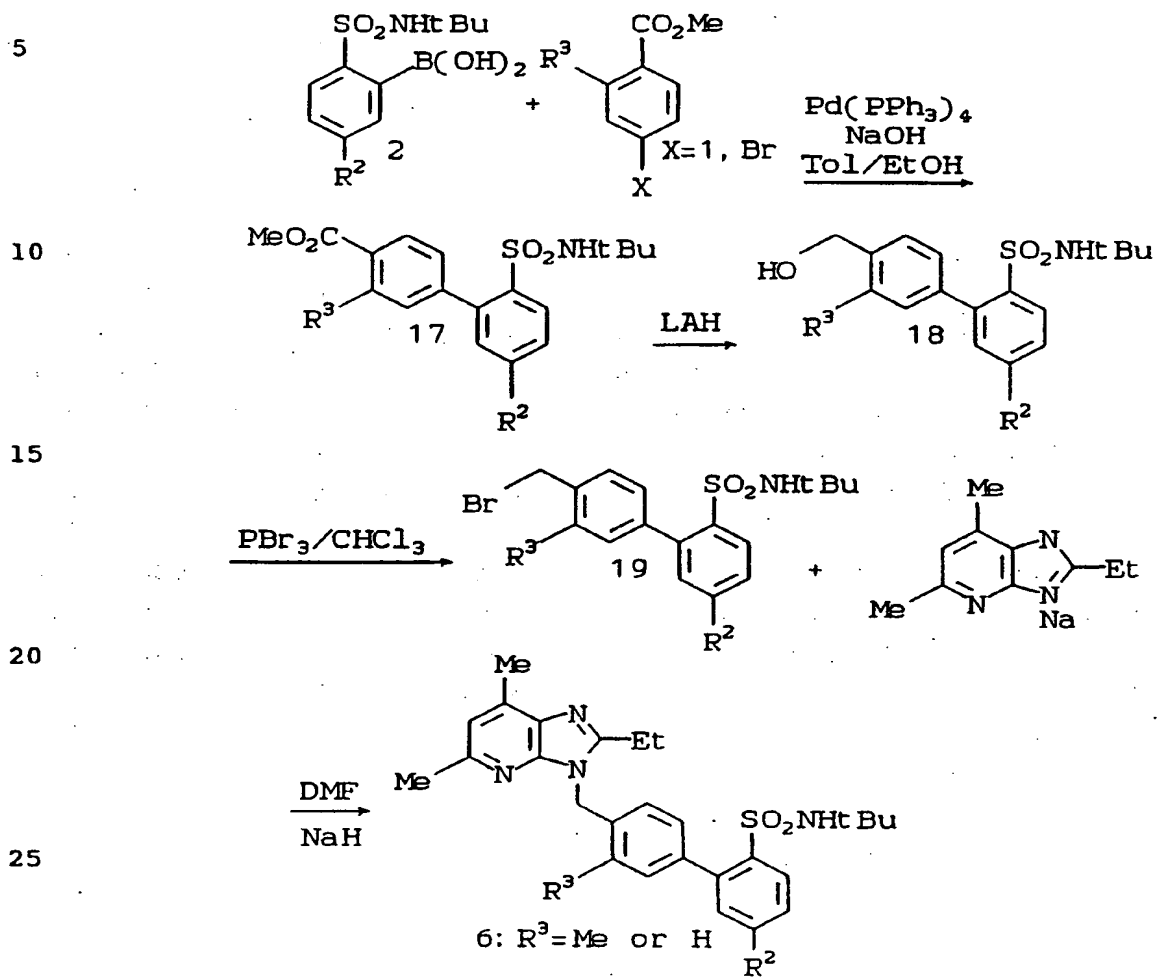
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SCHEME IV



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The compounds of this invention form salts with various inorganic and organic acids and bases which are also within the scope of the invention. Such salts include ammonium salts, alkali metal salts like sodium and potassium salts, alkaline earth metal salts like the calcium and magnesium salts, salts with organic bases; e.g., dicyclohexylamine salts, N-methyl-D-glucamine salts, salts with amino acids like arginine, lysine, and the like. Also, salts with organic and inorganic acids may be prepared; e.g., HCl, HBr, H₂SO₄, H₃PO₄, methanesulfonic, toluenesulfonic, maleic, fumaric, camphorsulfonic. The non-toxic, physiologically, acceptable salts are preferred, although other salts are also useful; e.g., in isolating or purifying the product.

The salts can be formed by conventional means such as by reacting the free acid or free base forms of the product with one or more equivalents of the appropriate base or acid in a solvent or medium in which the salt is insoluble, or in a solvent such as water which is then removed in vacuo or by freeze-drying or by exchanging the cations of an existing salt for another cation on a suitable ion exchange resin.

Angiotensin II (A II) is a powerful arterial vasoconstrictor, and it exerts its action by interacting with specific receptors present on cell membranes. The compounds described in the present invention act as competitive antagonists of A II at the receptors. In order to identify A II antagonists and determine their efficacy in vitro, the following three ligand-receptor binding assays were established.

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Receptor binding assay using rabbit aortae membrane preparation:

Three frozen rabbit aortae (obtained from Pel-Freeze Biologicals) were suspended in 5mM Tris-0.25M Sucrose, pH 7.4 buffer (50 ml) homogenized, and then centrifuged. The mixture was filtered through a cheesecloth and the supernatant was centrifuged for 30 minutes at 20,000 rpm at 4°C. The pellet thus obtained was resuspended in 30 ml of 50mM Tris-5 mM MgCl₂ buffer containing 0.2% Bovine Serum Albumin and 0.2 mg/ml Bacitracin and the suspension was used for 100 assay tubes. Samples tested for screening were done in duplicate. To the membrane preparation (0.25 ml) there was added ¹²⁵I-Sar¹Ile⁸-angiotensin II [obtained from New England Nuclear] (10μl; 20,000 cpm) with or without the test sample and the mixture was incubated at 37°C for 90 minutes. The mixture was then diluted with ice-cold 50mM Tris-0.9% NaCl, pH 7.4 (4ml) and filtered through a glass fiber filter (GF/B Whatman 2.4" diameter). The filter was soaked in scintillation cocktail (10 ml) and counted for radioactivity using Packard 2660 Tricarb liquid scintillation counter. The inhibitory concentration (IC₅₀) of potential A II antagonist which gives 50% displacement of the total specifically bound ¹²⁵I-Sar¹Ile⁸-angiotensin II was presented as a measure of the efficacy of such compounds as A II antagonists.

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Receptor assay using Bovine adrenal cortex preparation

Bovine adrenal cortex was selected as the source of A II receptor. Weighed tissue (0.1 g is needed for 100 assay tubes) was suspended in Tris.HCl (50mM), pH 7.7 buffer and homogenized. The homogenate was centrifuged at 20,000 rpm for 15 minutes. Supernatant was discarded and pellets resuspended in buffer [Na₂HPO₄ (10mM)-NaCl (120mM)-disodium EDTA (5mM) containing phenylmethane sulfonyl fluoride (PMSF)(0.1mM)]. (For screening of compounds, generally duplicates of tubes are used). To the membrane preparation (0.5 ml) there was added ³H-angiotensin II (50mM) (10μl) with or without the test sample and the mixture was incubated at 37°C for 1 hour. The mixture was then diluted with Tris buffer (4ml) and filtered through a glass fiber filter (GF/B Whatman 2.4" diameter). The filter was soaked in scintillation cocktail (10ml) and counted for radioactivity using Packard 2660 Tricarb liquid scintillation counter. The inhibitory concentration (IC₅₀) of potential A II antagonist which gives 50% displacement of the total specifically bound ³H-angiotensin II was presented as a measure of the efficacy of such compounds as A II antagonists.

Receptor assay using rat brain membrane preparation

Membranes from rat brain (thalamus, hypothalamus and midbrain) were prepared by homogenization in 50 mM Tris HCl (pH 7.4), and centrifuged at 50,000 x g. The resulting pellets were washed twice in 100 mM NaCl, 5 mM Na₂-EDTA, 10 mM Na₂HPO₄ (pH 7.4) and 0.1 mM PMSF by resuspension

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and centrifugation. For binding assays, the pellets were resuspended in 160 volumes of binding assay buffer (100 mM NaCl, 10 mM Na₂HPO₄, 5 mM Na₂•EDTA, pH 7.4, 0.1 mM PMSF, 0.2 mg/ml soybean trypsin inhibitor, 0.018 mg/ml o-phenanthroline, 77 mg/ml dithiothreitol and 0.14 mg/ml bacitracin. For ¹²⁵I-Ile⁸-angiotensin II binding assays, 10 µl of solvent (for total binding), Sar¹,Ile⁸-angiotensin II (1 µM) (for nonspecific binding) or test compounds (for displacement) and 10 µl of [¹²⁵I]Sar¹,Ile⁸-angiotensin II (23-46 pM) were added to duplicate tubes. The receptor membrane preparation (500 µl) was added to each tube to initiate the binding reaction. The reaction mixtures were incubated at 37°C for 90 minutes. The reaction was then terminated by filtration under reduced pressure through glass-fiber GF/B filters and washed immediately 4 times with 4 ml of 5 mM ice-cold Tris HCl (pH 7.6) containing 0.15 M NaCl. The radioactivity trapped on the filters was counted using a gamma counter.

The potential antihypertensive effects of the compounds described in the present invention may be evaluated using the methodology described below:

Male Charles River Sprague-Dawley rats (300-375 gm) were anesthetized with methohexital (Brevital; 50 mg/kg i.p.) and the trachea was cannulated with PE 205 tubing. A stainless steel pithing rod (1.5 mm thick, 150 mm long) was inserted into the orbit of the right eye and down the spinal column. The rats were immediately placed on a Harvard Rodent Ventilator (rate - 60 strokes per minute, volume -

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1.1 cc per 100 grams body weight). The right carotid artery was ligated, both left and right vagal nerves were cut, and the left carotid artery was cannulated with PE 50 tubing for drug administration, and body
5 temperature was maintained at 37°C by a thermostatically controlled heating pad which received input from a rectal temperature probe. Atropine (1 mg/kg i.v.) was then administered, and 15 minutes later propranolol (1 mg/kg i.v.). Thirty minutes later
10 angiotensin II or other agonists were administered intravenously at 30-minute intervals and the increase in the diastolic blood pressure was recorded before and after drug or vehicle administration.

Using the methodology described above,
15 representative compounds of the invention were evaluated and all were found to exhibit an activity of at least $IC_{50} 10\mu M$ against the AT_1 and AT_2 subtype receptors thereby demonstrating and confirming the utility of the compounds of the invention as
20 effective A II antagonists with "balanced" AT_1/AT_2 activity.

Thus, the compounds of the invention are useful in treating hypertension. They are also of value in the management of acute and chronic
25 congestive heart failure and angina. These compounds are also expected to be useful in primary and secondary hyperaldosteronism, renal diseases such as diabetic nephropathy, glomerulonephritis, glomerular sclerosis, nephrotic syndrome,
30 hypertensive nephrosclerosis, end stage renal disease, renal transplant therapy, renovascular hypertension, scleroderma, left ventricular

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dysfunction, systolic and diastolic dysfunction
diabetic retinopathy, in the management of vascular
disorders such as migraine or Raynaud's disease, as
prophylaxis to minimize the atherosclerotic process,
5 in neointimal hyperplasia following angioplasty or
vascular injury and to retard the onset of type II
diabetes. The application of the compounds of this
invention for these and similar disorders will be
apparent to those skilled in the art.

10 The compounds of this invention are also
useful to treat elevated intraocular pressure and to
enhance retinal blood flow and can be administered to
patients in need of such treatment with typical
pharmaceutical formulations such as tablets, capsules,
15 injectables and the like as well as topical ocular
formulations in the form of solutions, ointments,
inserts, gels, and the like. Pharmaceutical
formulations prepared to treat intraocular pressure
would typically contain about 0.1% to 15% by weight,
20 preferably 0.5% to 2% by weight, of a compound of
this invention. For this use, the compounds of this
invention may also be used in combination with other
medications for the treatment of glaucoma including
choline esterase inhibitors such as physostigmine
25 salicylate or demecarium bromide, parasympathomimetic
agents such as pilocarpine nitrate, β -adrenergic
antagonists such as timolol maleate, adrenergic
agonists such as epinephrine and carbonic anhydrase
inhibitors such as TRUSOPT™.

30 In the management of hypertension and the
clinical conditions noted above, the compounds of
this invention may be utilized in compositions such

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as tablets, capsules or elixirs for oral administration, suppositories for rectal administration, sterile solutions or suspensions for parenteral or intramuscular administration, and the like. The compounds of this invention can be administered to patients (animals and human) in need of such treatment in dosages that will provide optimal pharmaceutical efficacy. Although the dose will vary from patient to patient depending upon the nature and severity of disease, the patient's weight, special diets then being followed by a patient, concurrent medication, and other factors which those skilled in the art will recognize, the dosage range will generally be about 1 to 1000 mg. per patient per day which can be administered in single or multiple doses. Preferably, the dosage range will be about 5.0 to 500 mg. per patient per day; more preferably about 5 to 300 mg. per patient per day.

The compounds of this invention can also be administered in combination with other antihypertensives and/or diuretics. For example, the compounds of this invention can be given in combination with diuretics such as hydrochlorothiazide, chlorothiazide, chlorthalidone, methyclothiazide, furosemide, ethacrynic acid, triamterene, amiloride, atriopeptin and spironolactone; calcium channel blockers, such as diltiazem, felodipine, nifedipine, amlodipine, nimodipine, isradipine, nitrendipine and verapamil; β -adrenergic antagonists such as timolol, atenolol, metoprolol, propranolol, nadolol and pindolol; angiotensin converting enzyme inhibitors such as

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enalapril, lisinopril, captopril, ramipril, quinapril and zofenopril; renin inhibitors such as A-69729 and FK 906 and FK 744; α -adrenergic antagonists such as prazosin, doxazosin, and terazosin; sympatholytic
5 agents such as methyldopa, clonidine and guanabenz, atriopeptidase inhibitors (alone or with ANP) such as UK-79300; serotonin antagonists such as ketanserin; A_2 -adenosine receptor agonists such as CGS 22492C; potassium channel agonists such as pinacidil and
10 cromakalim; and various other antihypertensive drugs including reserpine, minoxidil, guanethidine, hydralazine hydrochloride and sodium nitroprusside as well as combinations of the above-named drugs.

Combinations useful in the management of
15 congestive heart failure include, in addition, compounds of this invention with cardiac stimulants such as dobutamine and xamoterol and phosphodiesterase inhibitors including amrinone and milrinone.

Typically, the individual daily dosages
20 for these combinations can range from about one-fifth of the minimally recommended clinical dosages to the maximum recommended levels for the entities when they are given singly.

To illustrate these combinations, one of the
25 angiotensin II antagonists of this invention effective clinically in the 5-500 milligrams per day range can be effectively combined at levels of the 1.0-500 milligrams per day range with the following compounds at the indicated per day dose range:
30 hydrochlorothiazide (6-100 mg), chlorothiazide (125-500 mg), ethacrynic acid (5-200 mg), amiloride (5-20 mg), furosemide (5-80 mg), propranolol (10-480

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mg), timolol maleate (1-20 mg), methyldopa (125-2000 mg), felodipine (1-20 mg), nifedipine (5-120 mg), nitrendipine (5-60 mg) and diltiazem (30-540 mg). In addition, triple drug combinations of hydrochlorothiazide (5-100 mg) plus amiloride (5-20 mg) plus an angiotensin II antagonist of this invention (1-500 mg) or hydrochlorothiazide (5-100 mg) plus timolol maleate (5-60) plus an angiotensin II antagonist of this invention (1-500 mg) or hydrochlorothiazide (5-200 mg) and nifedipine (5-60 mg) plus an angiotensin II antagonist of this invention (1-500 mg) are effective combinations to control blood pressure in hypertensive patients. Naturally, these dose ranges can be adjusted on a unit basis as necessary to permit divided daily dosage and, as noted above, the dose will vary depending on the nature and severity of the disease, weight of patient, special diets and other factors.

Typically, these combinations can be formulated into pharmaceutical compositions as discussed below.

About 1 to 100 mg. of compound or mixture of compounds of Formula I or a physiologically acceptable salt is compounded with a physiologically acceptable vehicle, carrier, excipient, binder, preservative, stabilizer, flavor, etc., in a unit dosage form as called for by accepted pharmaceutical practice. The amount of active substance in these compositions or preparations is such that a suitable dosage in the range indicated is obtained.

Illustrative of the adjuvants which can be incorporated in tablets, capsules and the like are

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the following: a binder such as gum tragacanth, acacia, corn starch or gelatin; an excipient such as microcrystalline cellulose; a disintegrating agent such as corn starch, pregelatinized starch, alginic acid and the like; a lubricant such as magnesium stearate; a sweetening agent such as sucrose, lactose or saccharin; a flavoring agent such as peppermint, oil of wintergreen or cherry. When the dosage unitform is a capsule, it may contain, in addition to materials of the above type, a liquid carrier such as fatty oil. Various other materials may be present as coatings or to otherwise modify the physical form of the dosage unit. For instance, tablets may be coated with shellac, sugar or both. A syrup or elixir may contain the active compound, sucrose as a sweetening agent, methyl and propyl parabens as preservatives, a dye and a flavoring such as cherry or orange flavor.

Sterile compositions for injection can be formulated according to conventional pharmaceutical practice by dissolving or suspending the active substance in a vehicle such as water for injection, a naturally occurring vegetable oil like sesame oil, coconut oil, peanut oil, cottonseed oil, etc., or a synthetic fatty vehicle like ethyl oleate or the like. Buffers, preservatives, antioxidants and the like can be incorporated as required.

The following examples illustrate the preparation of the compounds of formula (I) and their incorporation into pharmaceutical compositions and as such are not to be considered as limiting the invention set forth in the claims appended hereto.

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Example 1

5 5,7-dimethyl-2-ethyl-3[[2'-(N-n-pentylcarbonylsulfon-
amido)-5'-n-propyl[1,1'biphenyl]-4-yl]methylimidazo
[4,5-b]pyridine (compound 3 of Table 1)

Step A: Preparation of 4-n-propylbenzene-t-butyl-
sulfonamide (Scheme I, compound 2, R² =
n-pr)

10 To a solution of 4-n-propylphenylsulfonyl
chloride (Lancaster) in anhydrous CH₂Cl₂ (0.5 M
solution) cooled to 0°C under N₂ was added
t-butylamine (2.2 equiv) slowly through a dropping
15 funnel. After complete addition, the reaction was
stirred at rt for 12h. The CH₂Cl₂ was removed under
reduced pressure and the residue was extracted into
Et₂O and washed with 2N NaOH, H₂O and brine. The
organic was dried over anhydrous MgSO₄ and
concentrated in vacuo to afford the titled product.
20 R_f = 0.46 (3:1 Hex/EtOAc).
¹H NMR (200 MHz, CDCl₃) δ 0.93 (t, 3H), 1.22 (s, 9H),
1.62 (m, 2H), 2.65 (t, 2H), 4.67 (bs, 1H), 7.27 (d,
2H), 7.79 (d, 2H).

25 Step B: Preparation of 2-t-butylsulfonamido-5-n-
propylphenylboric acid (Scheme I, compound
3, R² = n-pr)

30 To a solution of 4-n-propylphenyl-t-butyl-
sulfonamide (2.85 g, 11.2 mmol) in anhydrous THF (20
mL) cooled to -40°C under N₂ was added 2.5M n-BuLi
solution (11.2 mL, 2.5 equiv). The mixture was
warmed to rt and stirred for 2h. To the mixture,
containing the bright red dianion at 0°C, was added

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B(OiPr)₃ (3.9 mL, 1.5 equiv). The next day 2N HCl (3 mL) was added and the mixture was stirred for 1h. The solvent was removed under reduced pressure and the residue was extracted with EtOAc. The organic was washed with 2N HCl, H₂O and brine. The organic was dried over anhydrous MgSO₄ and concentrated in vacuo to afford the titled compound. R_f = 0.5 (1:1 EtOAc/Hex). The crude material was used in subsequent steps without further purification.

Step C: Preparation of 5,7-dimethyl-2-ethyl-3[[2'-(N-tbutylsulfonamido)-5'-n-propyl-[1,1'biphenyl]-4-yl]methylimidazo[4,5-b]pyridine (Scheme I, compound 5, R² = n-pr)

To a solution of 5,7-dimethyl-2-ethyl-3[[4-bromo]phenyl]methylimidazo[4,5-b]pyridine (6.0 g, 17.4 mmol) and the product of step B (11.2 g, 37.3 mmol) in toluene (230 mL) was added 1.25 N NaOH (58 mL), EtOH (160 mL) and Pd(PPh₃)₄ (1.25 g, 3 mol %). The reaction mixture was stirred at 100° C under N₂ for 2 h. The solvent was removed under reduced pressure and the residue was taken up in EtOAc. The organic was washed with 1 N NaOH, H₂O and brine and dried over anhydrous MgSO₄ and concentrated in vacuo. The titled product was recrystallized from EtOAc/Hex. R_f = 0.5 (2:1 EtOAc/Hex).

¹H NMR (400 MHz, CD₃OD) δ 0.93 (t, 3H), 0.95 (s, 9H), 1.32 (t, 3H), 1.67 (m, 2H), 2.58 (s, 3H), 2.61 (s, 3H), 2.66 (t, 2H), 2.91 (q, 2H), 5.61 (s, 2H), 7.03 (s, 1H), 7.09 (d, 1H), 7.18 (d, 2H), 7.32 (dd, 1H), 7.41 (d, 2H), 7.97 (d, 1H).

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Step D: Preparation of 5,7-dimethyl-2-ethyl-3[[2'-(sulfonamido)-5'-n-propyl[1,1'-bi-phenyl]-4-yl]methylimidazo[4,5-b]pyridine (Scheme I, compound 5, $R^2 = n\text{-pr}$)

5 To a mixture of the product of step C (945 mg, 1.82 mmol) and anisole (0.5 mL) was added TFA (5 mL). After standing at rt for 24 h, the mixture was concentrated in vacuo. The residue was taken up in EtOAc and washed with 2N Na_2CO_3 solution, H_2O and
10 brine. The organic was dried over anhydrous MgSO_4 and concentrated in vacuo. The titled product, crystallized from Hex/ Et_2O , was obtained as a white powder. $R_f = 0.29$ (2:1 EtOAc/Hex).

15 Step E: Preparation of 5,7-dimethyl-2-ethyl-3[[2'-(N-n-pentylcarbonylsulfonamido)-5'-n-propyl[1,1'-biphenyl]-4-yl]methylimidazo[4,5-b]pyridine (compound 3 of Table 1)

To a solution of hexanoic acid (23 mg, 0.195 mmol) in dry THF (0.5 mL) under N_2 was added CDI (35
20 mg, 0.22 mmol). The mixture was stirred at 40° C for 2.5 h. To that solution was added a solution of the product of step D (30 mg, 0.065 mmol) and DBU (0.029 mL, 0.195 mmol) in THF (0.5 mL). The reaction was
25 stirred at 40° C for ca. 4 h. The reaction was quenched with MeOH (0.25 mL) and stirred for an additional 30 min. The solvent was removed and the the residue was dissolved in EtOAc and washed with 10% citric acid solution, H_2O and brine. The titled
30 product was purified by flash chromatography eluting with 80:10 :1 ($\text{CH}_2\text{Cl}_2/\text{MeOH}/\text{NH}_4\text{OH}$). $R_f = 0.37$ (20:1 $\text{CHCl}_3/\text{MeOH}$).

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¹H NMR (400 MHz, CD₃OD) δ 0.80 (t, 3H), 0.94 (t, 3H), 1.03 (m, 2H), 1.15 (m, 2H), 1.31 (m, 2H), 1.35 (t, 3H), 1.65 (m, 2H), 1.71 (t, 2H), 2.58 (s, 3H), 2.61 (s, 3H), 2.93 (m, 2H), 5.62 (s, 2H), 7.02 (s, 1H), 7.08 (d, 1H), 7.14 (d, 2H), 7.28 (d, 2H), 7.38 (dd, 1H), 8.05 (d, 1H).

Example 2

10 5,7-dimethyl-2-ethyl-3[[2'-(N-n-butylaminocarbonyl-sulfonamido)-5'-n-propyl-[1,1'biphenyl]-4-yl]methyl-imidazo[4,5-b]pyridine (compound 63 of Table 1)

To a solution of the product of Example 1, step D (75 mg, 0.162 mmol) in dry THF (2 mL) was added DBU (0.048 mL, 2 equiv) and n-butylisocyanate (0.182 mL, 10 equiv). After stirring at rt for 24 h, the solvent was removed under reduced pressure and the residue was dissolved in EtOAc and washed in 10% citric acid solution, H₂O and brine. The organic was dried over anhydrous MgSO₄ and concentrated in vacuo. The titled compound was purified by flash chromatography eluting with 80:10:1 (CH₂Cl₂/MeOH/NH₄OH). R_f = 0.68 (40:10:1 CHCl₃/MeOH/NH₄OH).

25 ¹H NMR (400 MHz, CD₃OD) δ 0.80 (t, 3H), 0.94 (t, 3H), 1.13 (m, 2H), 1.23 (m, 2H), 1.31 (t, 3H), 1.67 (m, 2H), 2.58 (s, 3H), 2.62 (s, 3H), 2.65 (t, 2H), 2.91 (m, 4H), 5.61 (s, 2H), 7.02 (s, 1H), 7.08 (s, 1H), 7.12 (d, 2H), 7.28 (d, 2H), 7.37 (d, 1H), 8.01 (d, 1H).

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Example 3

5 5,7-dimethyl-2-ethyl-3[[2'-(N-(2-phenylethyl)carbonyl-
sulfonamido)-5'-n-propyl[1,1'biphenyl]-4-yl]methyl-
imidazo[4,5-b]pyridine (compound 83 of Table 1)

To a solution of hydrocinnamic acid (50 mg, 0.33 mmol) in dry THF (1 mL) was added CDI (59 mg, 0.36 mmol). The mixture was stirred at 50° C for 2 h.
10 To that mixture was added a solution of the product of Example 1, step D (50 mg, 0.108 mmol) and DBU (0.050 mL, 0.33 mmol) in dry THF (1 mL). The reaction was stirred at 50° C for 12 h then quenched with MeOH (0.25 mL) and concentrated in vacuo. The
15 residue was dissolved in EtOAc and washed with 10% citric acid solution, H₂O and brine. The organic wqs dried over anhydrous MgSO₄ and concentrated in vacuo. The titled product was purified by radial chromatography eluting with 100:10:1
20 (CH₂Cl₂/MeOH/NH₄OH). R_f = 0.56 (80:10:1 CHCl₃/MeOH/NH₄OH).
¹H NMR (400 MHz, CD₃OD) δ 0.94 (t, 3H), 1.28 (t, 3H), 1.62 (m, 2H), 2.13 (t, 2H), 2.55 (s, 3H), 2.62 (s, 3H), 2.85 (q, 2H), 5.55 (s, 2H), 6.95-7.03 (comp m, 6H), 7.08 (m, 3H), 7.13 (d, 2H), 7.31 (dd, 1H), 8.02 (d, 1H).
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Example 4

5 5,7-dimethyl-2-ethyl-3[[2'-(N-(2-phenoxyphenyl)car-
bonylsulfonamido)-5'-n-propyl[1,1'biphenyl]-4-yl]
methylimidazo[4,5-b]pyridine (compound 103 of Table 1)

To a solution of 2-phenoxybenzoic acid acid
(138 mg, 0.644 mmol) in dry THF (2 mL) was added CDI
(125 mg, 0.71 mmol). The mixture was stirred at 40° C
10 for 2.5 h. To that mixture was added a solution of
the product of Example 1, step D (100 mg, 0.216 mmol)
and DBU (0.10 mL, 0.66 mmol) in dry THF (2 mL). The
reaction was stirred at 40° C for 3.5 h then quenched
with MeOH (0.25 mL) and concentrated in vacuo. The
15 residue was dissolved in EtOAc and washed with 10%
citric acid solution, H₂O and brine. The organic wqs
dried over anhydrous MgSO₄ and concentrated in
vacuo. The titled product was purified by radial
chromatography eluting with 100:7:1
20 (CH₂Cl₂/MeOH/NH₄OH). R_f = 0.42 (2:1 EtOAc/Hex).
¹H NMR (400 MHz, CD₃OD) δ 0.91 (t, 3H), 1.26 (t, 3H),
1.62 (m, 2H), 2.55 (s, 3H), 2.61 (s, 3H), 2.71 (q,
2H), 5.50 (s, 2H), 6.72 (d, 1H), 6.95-7.03 (comp m,
6H), 7.21 (comp m, 6H), 7.43 (d, 1H), 8.12 (d, 1H).

25

Example 5

5,7-dimethyl-2-ethyl-3[[2'-(N-benzenecarbonylsulfon-
amido)-5'-n-propyl[1,1'biphenyl]-4-yl]methylimidazo
30 [4,5-b]pyridine (compound 143 of Table 1)

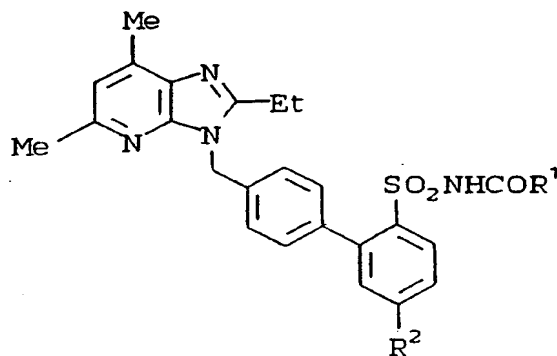
- 44 -

To a solution of the product from Example 1, step D (100 mg, 0.216 mmol) in dry pyridine (2 mL) was added DMAP (20 mg) and benzoyl chloride (300 mg, 10 equiv). After stirring for 6 h the reaction was quenched with MeOH (0.5 mL) and the solvent was removed in vacuo. The residue was taken up in EtOAc and washed with 10% citric acid, H₂O and brine. The titled product was purified by flash chromatography eluting with 60:10:1 (CH₂Cl₂/MeOH/NH₄OH). R_f = 0.56 (80:10:1 CHCl₃/MeOH/NH₄OH).

¹H NMR (200 MHz, CDCl₃) δ 0.83 (t, 3H), 1.29 (t, 3H), 1.58 (m, 2H), 2.53 (s, 3H), 2.61 (s, 3H), 2.84 (q, 2H), 5.48 (s, 2H), 6.85–7.01 (comp m, 4H), 7.09–7.29 (comp m, 6H), 7.38 (d, 2H), 8.17 (d, 1H).

Examples 6 through 13, shown in Table 2, were prepared using procedures described in the previous five examples and illustrated in Schemes I through IV.

TABLE 2



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EXAMPLE	R ¹	R ²
6	(CH ₂) ₄ CH ₃	CH ₂ N(CH ₃) ₂
7	NHn-Bu	O-n-Bu
5 8	(CH ₂) ₄ CH ₃	CH(CH ₃) ₂
9	CH ₂ OEt	(CH ₂) ₂ CH ₃
10 10	CH ₂ On-Bu	(CH ₂) ₂ CH ₃
11	(CH ₂) ₄ CH ₃	CH ₂ N(CH ₂ CH ₂) ₂
12	(CH ₂) ₄ CH ₃	CH ₂ N(CH ₂ CH ₂) ₂ O
10 13	(CH ₂) ₄ CH ₃	O-n-Bu

Example 6

15 5,7-dimethyl-2-ethyl-3[[2'-(N-n-pentylcarbonylsulfon-
amido)-5'-dimethylaminomethyl [1,1'biphenyl]-4-yl]
methylimidazo[4,5-b]pyridine (compound 15 of Table 1)

The titled compound was prepared as follows: Intermediate 2 of scheme I (R² =
20 CH₂N(CH₃)₂) was prepared using the method described in scheme II D. Completion of the antagonist was carried out as illustrated in scheme I. The final step, preparation of compound 7 of scheme I (R¹ =
25 (CH₂)₄CH₃, R² = CH₂N(CH₃)₂) from the free sulfonamide, was carried out using hexanoic acid and CDI (method A of scheme I).

¹H NMR (400 MHz, CD₃OD) δ 0.81 (t, 3H), 1.08 (m, 2H),
1.18 (m, 2H), 1.31 (m, 2H), 1.34 (t, 3H), 1.78 (t,
2H), 2.49 (s, 6H), 2.58 (s, 3H), 2.61 (s, 3H), 2.91
30 (q, 2H), 3.84 (s, 2H), 5.59 (s, 2H), 7.01 (s, 1H),
7.12 (d, 2H), 7.23 (d, 1H), 7.32 (d, 2H), 7.48 (dd,
1H), 8.12 (d, 1H).

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Example 7

5 5,7-dimethyl-2-ethyl-3[[2'-(N-n-butylaminocarbonyl-sulfonamido)-5'-n-butoxy[1,1'biphenyl]-4-yl]methyl-imidazo[4,5-b]pyridine(compound 66 of Table 1)

The titled compound was prepared as follows: Intermediate 2 of scheme I ($R^2 = O(CH_2)_3CH_3$) was prepared from commercially available 4-butoxybenzenesulfonyl chloride. Completion of the antagonist was carried out as illustrated in scheme I. The final step, preparation of compound 7 of scheme I ($R^1 = NH(CH_2)_3CH_3$, $R^2 = O(CH_2)_3CH_3$) from the free sulfonamide, was carried out using n-butylisocyanate and DBU as base (method C of scheme I).
15 1H NMR (400 MHz, CD_3OD) δ 0.81 (t, 3H), 0.96 (t, 3H), 1.13 (m, 2H), 1.23 (m, 2H), 1.33 (t, 3H), 1.47 (m, 2H), 1.75 (m, 2H), 2.58 (s, 3H), 2.61 (s, 3H), 2.91 (q, 4H), 4.03 (t, 2H), 5.60 (s, 2H), 6.73 (d, 1H),
20 7.01 (s, 1H), 7.02 (dd, 1H), 7.13 (d, 2H), 7.28 (d, 2H), 8.03 (d, 1H).

Example 8

25 5,7-dimethyl-2-ethyl-3[[2'-(N-n-pentylcarbonylsulfon-amido)-5'-isopropyl[1,1'biphenyl]-4-yl]methylimidazo[4,5-b]pyridine (compound 5 of Table 1)

The titled compound was prepared as follows: Intermediate 2 of scheme I ($R^2 = CH(CH_3)_2$) was prepared from commercially available 4-butoxybenzenesulfonyl chloride. Completion of the

30

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antagonist was carried out as illustrated in scheme I. The final step, preparation of compound 7 of scheme I ($R^1 = (CH_2)_4CH_3$, $R^2 = CH(CH_3)_2$) from the free sulfonamide, was carried out using hexanoic acid and CDI (method A of scheme I).

¹H NMR (400 MHz, CD₃OD) δ 0.79 (t, 3H), 1.05 (m, 2H), 1.17 (m, 2H), 1.24 (d, 6H), 1.33 (t, 3H), 1.34 (m, 2H), 1.79 (t, 2H), 2.57 (s, 3H), 2.61 (s, 3H), 2.91 (m, 3H), 5.61 (s, 2H), 7.01 (s, 1H), 7.08 (d, 1H), 7.13 (d, 2H), 7.28 (d, 2H), 7.40 (dd, 1H), 8.07 (d, 1H).

Example 9

5,7-dimethyl-2-ethyl-3[[2'-(N-ethoxymethylcarbonylsulfonamido)-5'-n-propyl[1,1'biphenyl]-4-yl]methylimidazo[4,5-b]pyridine (compound 43 of Table 1)

The titled compound was prepared from the product of Example 1, step D. Completion of the antagonist was carried out as illustrated in scheme I. The final step, preparation of compound 7 of scheme I ($R^1 = CH_2OCH_2CH_3$, $R^2 = (CH_2)_2CH_3$) from the free sulfonamide, was carried out using ethoxyacetic acid and CDI (method A of scheme I).

¹H NMR (400 MHz, CD₃OD) δ 0.93 (t, 3H), 1.04 (t, 3H), 1.35 (t, 3H), 1.62 (m, 2H), 2.58 (s, 3H), 2.60 (s, 3H), 2.61 (t, 2H), 2.91 (q, 2H), 3.28 (t, 2H), 3.45 (s, 2H), 5.59 (s, 2H), 6.99 (d, 1H), 7.01 (s, 1H), 7.11 (d, 2H), 7.28 (dd, 1H), 7.33 (d, 2H), 8.02 (d, 1H).

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Example 10

5 5,7-dimethyl-2-ethyl-3[[2'-(N-(n-butoxy)methylcar-
bonylsulfonamido)-5'-n-propyl[1.1'biphenyl]-4-yl]
methylimidazo[4,5-b]pyridine (compound 23 of Table 1)

The titled compound was prepared from the product of Example 1, step D. Completion of the antagonist was carried out as illustrated in scheme I. The final step, preparation of compound 7 of scheme I ($R^1 = \text{CH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$, $R^2 = (\text{CH}_2)_2\text{CH}_3$) from the free sulfonamide, was carried out using n-butoxyacetic acid and CDI (method A of scheme I). ^1H NMR (400 MHz, CD_3OD) δ 0.84 (t, 3H), 0.93 (t, 3H), 1.22 (m, 2H), 1.35 (t, 3H), 1.39 (m, 2H), 1.62 (m, 2H), 2.58 (s, 3H), 2.61 (s, 3H), 2.62 (t, 2H), 2.91 (q, 2H), 3.21 (t, 2H), 3.43 (s, 2H), 5.59 (s, 2H), 6.99 (d, 1H), 7.01 (s, 1H), 7.11 (d, 2H), 7.28 (dd, 1H), 7.32 (d, 2H), 8.03 (d, 1H).

Example 11

25 5,7-dimethyl-2-ethyl-3[[2'-(N-n-pentylcarbonylsulfon-
amido)-5'-pyrrolidin-1-ylmethyl[1.1'biphenyl]-4-yl]
methylimidazo[4,5-b]pyridine (compound 16 of Table 1)

The titled compound was prepared as follows: Intermediate 2 of scheme I ($R^2 = \text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2$) was prepared using the method described in scheme II D. Completion of the antagonist was carried out as illustrated in scheme I. The final step, preparation of compound 7 of

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scheme I ($R^1 = (CH_2)_4CH_3$, $R^2 = CH_2N(CH_2CH_2)_2$) from the free sulfonamide, was carried out using hexanoic acid and CDI (method A of scheme I).

¹H NMR (400 MHz, CD₃OD) δ 0.81 (t, 3H), 1.11 (m, 2H),
5 1.19 (m, 2H), 1.32 (m, 5H), 1.79 (t, 2H), 1.98 (bs,
4H), 2.57 (s, 3H), 2.60 (s, 3H), 2.91 (q, 2H), 3.12
(bs, 4H), 4.21 (s, 2H), 5.59 (s, 2H), 6.99 (s, 1H),
7.11 (d, 2H), 7.28 (d, 1H), 7.31 (d, 2H), 7.51 (dd,
1H), 8.10 (d, 1H).

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Example 12

5,7-dimethyl-2-ethyl-3[[2'-(N-n-pentylcarbonylsulfon-
amido)-5'-morpholin-1-ylmethyl[1,1'-biphenyl]-4-yl]
15 methylimidazo[4,5-b]pyridine (compound 17 of Table 1)

The titled compound was prepared as follows: Intermediate 2 of scheme I ($R^2 = CH_2N(CH_2CH_2)_2O$) was prepared using the method
20 described in scheme II D. Completion of the antagonist was carried out as illustrated in scheme I. The final step, preparation of compound 7 of scheme I ($R^1 = (CH_2)_4CH_3$, $R^2 = CH_2N(CH_2CH_2)_2O$) from the free sulfonamide, was carried out using hexanoic
25 acid and CDI (method A of scheme I).

¹H NMR (400 MHz, CD₃OD) δ 0.80 (t, 3H), 1.04 (m, 2H),
1.15 (m, 2H), 1.31-1.39 (m, 5H), 1.80 (t, 2H), 2.50
(bs, 4H), 2.58 (s, 3H), 2.62 (s, 3H), 2.92 (q, 2H),
3.62 (bs, 2H), 3.68 (bm, 4H), 5.61 (s, 2H), 7.04 (s,
30 1H), 7.15 (d, 2H), 7.26 (s, 1H), 7.27 (d, 2H), 7.56
(dd, 1H), 8.12 (d, 1H).

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Example 13

5,7-dimethyl-2-ethyl-3[[2'-(N-n-pentylcarbonylsulfon-
amido)-5'-n-butoxy[1,1'biphenyl]-4-yl]methyylimidazo
5 [4,5-b]pyridine (compound 6 of Table 1)

The titled compound was prepared as follows: Intermediate 2 of scheme I ($R^2 =$
10 $O(CH_2)_3CH_3$) was prepared from commercially available 4-butoxybenzenesulfonyl chloride. Completion of the antagonist was carried out as illustrated in scheme I. The final step, preparation of compound 7 of
15 scheme I ($R^1 = (CH_2)_4CH_3$, $R^2 = O(CH_2)_3CH_3$) from the free sulfonamide, was carried out using hexanoic acid and CDI (method A of scheme I).

1H NMR (400 MHz, CD_3OD) δ 0.80 (t, 3H), 0.96 (t, 3H),
1.07 (m, 2H), 1.17 (m, 2H), 1.32 (m, 2H), 1.35 (t,
3H), 1.49 (m, 2H), 1.75 (m, 2H), 1.83 (t, 2H), 2.58
(s, 3H), 2.61 (s, 3H), 2.92 (q, 4H), 4.04 (t, 2H),
20 5.61 (s, 2H), 6.73 (d, 1H), 7.02 (s, 1H), 7.04 (dd,
1H), 7.13 (d, 2H), 7.28 (d, 2H), 8.09 (d, 1H).

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FORMULATION EXAMPLESTypical Pharmaceutical Compositions Containing a
Compound of the Invention

5

A: Dry Filled Capsules Containing 50 mg of Active
Ingredient Per Capsule

	<u>Ingredient</u>	<u>Amount per Capsule (mg)</u>
10	Compound 1	50
	Lactose	149
	Magnesium stearate	<u>1</u>
	Capsule (size No. 1)	200

15 Compound 1 can be reduced to a No. 60 powder
and the lactose and magnesium stearate can then be
passed through a No. 60 blotting cloth onto the
powder. The combined ingredients can then be mixed
for about 10 minutes and filled into a No. 1 dry
20 gelatin capsule.

B: Tablet

25 A typical tablet would contain Compound 1
(25 mg), pregelatinized starch USP (82 mg), micro-
crystalline cellulose (82 mg) and magnesium stearate
(1 mg).

C: Combination Tablet

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A typical suppository formulations for rectal administration can contain Compound 1 (1-25 mg), butylated hydroxyanisole (0.08-1.0 mg), disodium calcium edetate (0.25-0.5 mg), and polyethylene glycol (775-1600 mg). Other suppository formulations can be made by substituting, for example, butylated hydroxytoluene (0.04-0.08 mg) for the disodium calcium edetate and a hydrogenated vegetable oil (675-1400 mg) such as Suppocire L, Wecobee FS, Wecobee M, Witepsols, and the like, for the polyethylene glucol. Further, these suppository formulations can also include another active ingredient such as another antihypertensive and/or a diuretic and/or an angiotensin converting enzyme and/or a calcium channel blocker in pharmaceutically effective amounts as described, for example, in C above.

E: Injection

20

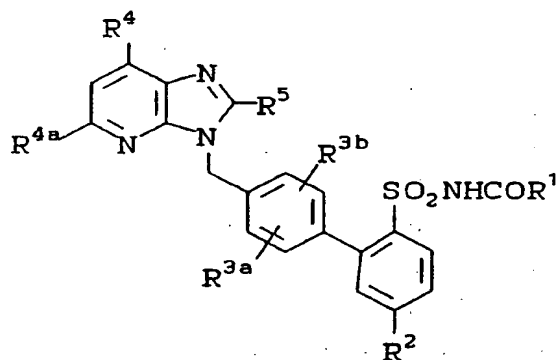
A typical injectable formulation would contain Compound 1 (5.42 mg), sodium phosphate dibasic anhydrous (11.4 mg) benzyl alcohol (0.01 ml) and water for injection (1.0 ml). Such an injectable formulation can also include a pharmaceutically effective amount of another active ingredient such as another antihypertensive and/or a diuretic and/or an angiotensin converting enzyme inhibitor and/or a calcium channel blocker.

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WHAT IS CLAIMED IS:

1. A compound of structural formula:



or a pharmaceutically acceptable salt thereof, wherein

- R^1 is
- C_{1-6} alkyl,
 - C_{1-6} alkylamino,
 - C_{1-6} alkoxy- $(CH_2)_n$ -, wherein n is 1 or 2,
 - aryl $S(O)_q$ -, wherein q is 0 to 3,
 - C_{1-6} alkylthio- $(CH_2)_n$ -,
 - aryl, either unsubstituted or substituted with
 - C_{1-6} alkyl,
 - aryloxy,
 - C_{1-6} alkoxy,
 - Cl,
 - Br, or
 - C_{1-6} alkylamino;

- R^2 is
- Cl,
 - C_{1-6} alkyl,

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- 5 c) C₁₋₅ alkoxy,
d) C₁₋₅ alkoxy-CH₂-,
e) di(C₁₋₅ alkyl) amino-CH₂-,
f) pyrrolidin-1-yl-CH₂-,
g) morpholin-1-yl-CH₂-,
h) polyfluoro-C₁₋₅ alkoxy,
i) aryl,
j) C₁₋₅ alkyl S(O)_q-(CH₂)_q,
k) aryl-(CH₂)_n-;

10

R^{3a} and R^{3b} are independently

- a) H,
b) F, Cl, Br or I,
c) C₁₋₄ alkyl,
15 d) C₁₋₄ alkoxy, or
e) aryl;

R^{3a} and R^{3b} on adjacent carbons can be joined together to form a benzo group;

20

R⁴ and R^{4a} are independently

- a) C₁₋₃ alkyl,
b) polyfluoro-C₁₋₃ alkyl,
25 c) -COHNR¹,
d) -CO₂R¹ or
e) -CONH (CH₂)_n-aryl; and

R⁵ is hydrogen or C₁₋₅ alkyl;

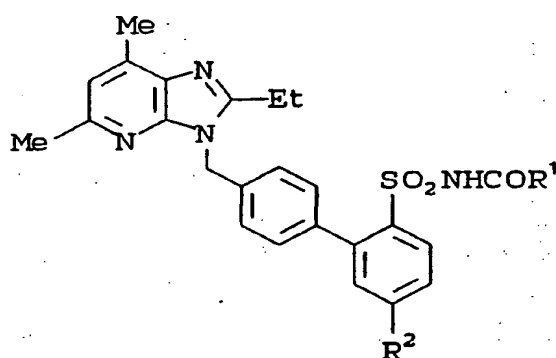
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2. The compound of Claim 1, or a pharmaceutically acceptable salt thereof, wherein R⁴ and R^{4a} are both C₁₋₃ alkyl, and R⁵ is C₁₋₅ alkyl.

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3. The compound of Claim 2, or a pharmaceutically acceptable salt thereof, wherein R^4 and R^{4a} are both methyl and R^5 is ethyl.

5 4. The compound of Claim 3, or a pharmaceutically acceptable salt thereof selected from the group comprising those in the following list:



20 $\text{SO}_2\text{NHCOR}^1$

R^2

$\text{SO}_2\text{NHCO}(\text{CH}_2)_4\text{CH}_3$

CH_3

$\text{SO}_2\text{NHCO}(\text{CH}_2)_4\text{CH}_3$

CH_2CH_3

25 $\text{SO}_2\text{NHCO}(\text{CH}_2)_4\text{CH}_3$

$(\text{CH}_2)_2\text{CH}_3$

$\text{SO}_2\text{NHCO}(\text{CH}_2)_4\text{CH}_3$

$(\text{CH}_2)_3\text{CH}_3$

$\text{SO}_2\text{NHCO}(\text{CH}_2)_4\text{CH}_3$

$\text{CH}(\text{CH}_3)_2$

$\text{SO}_2\text{NHCO}(\text{CH}_2)_4\text{CH}_3$

$\text{O}(\text{CH}_2)_3\text{CH}_3$

$\text{SO}_2\text{NHCO}(\text{CH}_2)_4\text{CH}_3$

$\text{CH}_2\text{CH}(\text{CH}_3)_2$

30 $\text{SO}_2\text{NHCO}(\text{CH}_2)_4\text{CH}_3$

OCH_3

$\text{SO}_2\text{NHCO}(\text{CH}_2)_4\text{CH}_3$

CH_2SCH_3

$\text{SO}_2\text{NHCO}(\text{CH}_2)_4\text{CH}_3$

CH_2OCH_3

$\text{SO}_2\text{NHCO}(\text{CH}_2)_4\text{CH}_3$

OCH_2CH_3

$\text{SO}_2\text{NHCO}(\text{CH}_2)_4\text{CH}_3$

Ph

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	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{C}(\text{CH}_3)_2\text{CH}_2\text{CH}_3$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{C}(\text{CH}_3)_3$
5	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2\text{O}$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	OCH_2CF_3
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{OCH}(\text{CH}_3)_2$
10	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	SCH_2CH_3
	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	CH_3
	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	CH_2CH_3
	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$(\text{CH}_2)_2\text{CH}_3$
	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$(\text{CH}_2)_3\text{CH}_3$
15	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{O}(\text{CH}_2)_3\text{CH}_3$
	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{CH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	OCH_3
	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	CH_2SCH_3
20	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	CH_2OCH_3
	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	OCH_2CH_3
	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	Ph
	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{C}(\text{CH}_3)_2\text{CH}_2\text{CH}_3$
	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{C}(\text{CH}_3)_3$
25	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2$
	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2\text{O}$
	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	OCH_2CF_3
	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	$\text{OCH}(\text{CH}_3)_2$
30	$\text{SO}_2\text{NHCOCCH}_2\text{O}(\text{CH}_2)_3\text{CH}_3$	SCH_2CH_3
	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	CH_3
	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	CH_2CH_3
	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	$(\text{CH}_2)_2\text{CH}_3$

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	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	$(\text{CH}_2)_3\text{CH}_3$
	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	$\text{CH}(\text{CH}_3)_2$
5	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	$\text{O}(\text{CH}_2)_3\text{CH}_3$
	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	$\text{CH}_2\text{CH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	OCH_3
	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	CH_2SCH_3
	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	CH_2OCH_3
10	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	OCH_2CH_3
	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	Ph
	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	$\text{C}(\text{CH}_3)_2\text{CH}_2\text{CH}_3$
	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	$\text{C}(\text{CH}_3)_3$
	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_3)_2$
15	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2$
	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2\text{O}$
	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	OCH_2CF_3
	$\text{SO}_2\text{NHCOCCH}_2\text{OCH}_2\text{CH}_3$	$\text{OCH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	SCH_2CH_3
20	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	CH_3
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	CH_2CH_3
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	$(\text{CH}_2)_2\text{CH}_3$
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	$(\text{CH}_2)_3\text{CH}_3$
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}(\text{CH}_3)_2$
25	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	$\text{O}(\text{CH}_2)_3\text{CH}_3$
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{CH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	OCH_3
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	CH_2SCH_3
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	CH_2OCH_3
30	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	OCH_2CH_3
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	Ph
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	$\text{C}(\text{CH}_3)_2\text{CH}_2\text{CH}_3$

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	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	$\text{C}(\text{CH}_3)_3$
	$\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_3)_2$
5	$\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2$
	$\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2\text{O}$
	$\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	OCH_2CF_3
	$\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	$\text{OCH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	SCH_2CH_3
10	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	CH_3
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	CH_2CH_3
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$(\text{CH}_2)_2\text{CH}_3$
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$(\text{CH}_2)_3\text{CH}_3$
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{CH}(\text{CH}_3)_2$
15	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{O}(\text{CH}_2)_3\text{CH}_3$
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{CH}_2\text{CH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	OCH_3
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	CH_2SCH_3
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	CH_2OCH_3
20	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	OCH_2CH_3
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	Ph
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{C}(\text{CH}_3)_2\text{CH}_2\text{CH}_3$
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{C}(\text{CH}_3)_3$
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{CH}_2\text{N}(\text{CH}_3)_2$
25	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2$
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2\text{O}$
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	OCH_2CF_3
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{OCH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	SCH_2CH_3
30	$\text{SO}_2\text{NHCO}(2\text{-PhO})\text{Ph}$	CH_3
	$\text{SO}_2\text{NHCO}(2\text{-PhO})\text{Ph}$	CH_2CH_3
	$\text{SO}_2\text{NHCO}(2\text{-PhO})\text{Ph}$	$(\text{CH}_2)_2\text{CH}_3$

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	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
5	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
10	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
15	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
20	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
25	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
30	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2

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	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
5	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
10	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
15	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
20	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
25	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
30	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2

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	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{CH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{O}(\text{CH}_2)_3\text{CH}_3$
5	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	OCH_3
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	CH_2SCH_3
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	CH_2OCH_3
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	OCH_2CH_3
10	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	Ph
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{C}(\text{CH}_3)_2\text{CH}_2\text{CH}_3$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{C}(\text{CH}_3)_3$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2$
15	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2\text{O}$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	OCH_2CF_3
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{OCH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	SCH_2CH_3
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	CH_3
20	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	CH_2CH_3
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$(\text{CH}_2)_2\text{CH}_3$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$(\text{CH}_2)_2\text{CH}_3$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{CH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{O}(\text{CH}_2)_3\text{CH}_3$
25	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	OCH_3
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	CH_2SCH_3
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	CH_2OCH_3
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	OCH_2CH_3
30	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	Ph
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{C}(\text{CH}_3)_2\text{CH}_2\text{CH}_3$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{C}(\text{CH}_3)_3$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_4\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_3)_2$

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	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
5	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
10	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
15	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
20	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
25	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
30	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOR}^1$	R^2

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	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{CH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	OCH_3
5	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	CH_2SCH_3
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	CH_2OCH_3
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	OCH_2CH_3
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	Ph
	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	$\text{C}(\text{CH}_3)_2\text{CH}_2\text{CH}_3$
10	$\text{SO}_2\text{NHCONH}(\text{CH}_2)_3\text{CH}_3$	$\text{C}(\text{CH}_3)_3$
	$\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2$
	$\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2\text{O}$
	$\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	OCH_2CF_3
15	$\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	$\text{OCH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCON}(\text{CH}_2)_3\text{CH}_3$	SCH_2CH_3
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	CH_3
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	CH_2CH_3
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$(\text{CH}_2)_2\text{CH}_3$
20	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$(\text{CH}_2)_3\text{CH}_3$
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{CH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{O}(\text{CH}_2)_3\text{CH}_3$
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{CH}_2\text{CH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	OCH_3
25	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	CH_2SCH_3
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	CH_2OCH_3
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	OCH_2CH_3
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	Ph
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{C}(\text{CH}_3)_2\text{CH}_2\text{CH}_3$
30	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{C}(\text{CH}_3)_3$
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{CH}_2\text{N}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCO}(\text{CH}_2)_2\text{Ph}$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2$

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	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_2\text{Ph}$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2\text{O}$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_2\text{Ph}$	OCH_2CF_3
5	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_2\text{Ph}$	$\text{OCH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOC}(\text{CH}_2)_2\text{Ph}$	SCH_2CH_3
	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	CH_3
	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	CH_2CH_3
	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	$(\text{CH}_2)_2\text{CH}_3$
10	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	$(\text{CH}_2)_2\text{CH}_3$
	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	$\text{CH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	$\text{O}(\text{CH}_2)_3\text{CH}_3$
	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	$\text{CH}_2\text{CH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	OCH_3
15	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	CH_2SCH_3
	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	CH_2OCH_3
	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	OCH_2CH_3
	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	Ph
	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	$\text{C}(\text{CH}_3)_2\text{CH}_2\text{CH}_3$
20	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	$\text{C}(\text{CH}_3)_3$
	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	$\text{CH}_2\text{N}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2$
	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	$\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2)_2\text{O}$
	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	OCH_2CF_3
25	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	$\text{OCH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOC}(2\text{-PhO})\text{Ph}$	SCH_2CH_3
	$\text{SO}_2\text{NHCOC}(2\text{-EtO})\text{Ph}$	CH_3
	$\text{SO}_2\text{NHCOC}(2\text{-EtO})\text{Ph}$	CH_2CH_3
	$\text{SO}_2\text{NHCOC}(2\text{-EtO})\text{Ph}$	$(\text{CH}_2)_2\text{CH}_3$
30	$\text{SO}_2\text{NHCOC}(2\text{-EtO})\text{Ph}$	$(\text{CH}_2)_2\text{CH}_3$
	$\text{SO}_2\text{NHCOC}(2\text{-EtO})\text{Ph}$	$\text{CH}(\text{CH}_3)_2$
	$\text{SO}_2\text{NHCOC}(2\text{-EtO})\text{Ph}$	$\text{O}(\text{CH}_2)_3\text{CH}_3$

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	$\text{SO}_2\text{NHCOR}^1$	R^2
	$\text{SO}_2\text{NHCOC(2-EtO)Ph}$	$\text{CH}_2\text{CH(CH}_3)_2$
	$\text{SO}_2\text{NHCOC(2-EtO)Ph}$	OCH_3
5	$\text{SO}_2\text{NHCOC(2-EtO)Ph}$	CH_2SCH_3
	$\text{SO}_2\text{NHCOC(2-EtO)Ph}$	CH_2OCH_3
	$\text{SO}_2\text{NHCOC(2-EtO)Ph}$	OCH_2CH_3
	$\text{SO}_2\text{NHCOC(2-EtO)Ph}$	Ph
	$\text{SO}_2\text{NHCOC(2-EtO)Ph}$	$\text{C(CH}_3)_2\text{CH}_2\text{CH}_3$
10	$\text{SO}_2\text{NHCOC(2-EtO)Ph}$	$\text{C(CH}_3)_3$
	$\text{SO}_2\text{NHCOC(2-EtO)Ph}$	$\text{CH}_2\text{N(CH}_3)_2$
	$\text{SO}_2\text{NHCOC(2-EtO)Ph}$	$\text{CH}_2\text{N(CH}_2\text{CH}_2)_2$
	$\text{SO}_2\text{NHCOC(2-PhO)Ph}$	OCH_2CF_3
	$\text{SO}_2\text{NHCOC(2-PhO)Ph}$	$\text{OCH(CH}_3)_2$
15	$\text{SO}_2\text{NHCOC(2-PhO)Ph}$	SCH_2CH_3
	SO_2NHCOPh	CH_3
	SO_2NHCOPh	CH_2CH_3
	SO_2NHCOPh	$(\text{CH}_2)_2\text{CH}_3$
	SO_2NHCOPh	$(\text{CH}_2)_2\text{CH}_3$
20	SO_2NHCOPh	$\text{CH(CH}_3)_2$
	SO_2NHCOPh	$\text{O(CH}_2)_3\text{CH}_3$
	SO_2NHCOPh	$\text{CH}_2\text{N(CH}_3)_2$
	SO_2NHCOPh	OCH_3
	SO_2NHCOPh	CH_2SCH_3
25	SO_2NHCOPh	CH_2OCH_3
	SO_2NHCOPh	OCH_2CH_3
	SO_2NHCOPh	Ph
	SO_2NHCOPh	$\text{C(CH}_3)_2\text{CH}_2\text{CH}_3$
	SO_2NHCOPh	$\text{C(CH}_3)_3$
30	SO_2NHCOPh	$\text{CH}_2\text{N(CH}_3)_2$
	SO_2NHCOPh	$\text{CH}_2\text{N(CH}_2\text{CH}_2)_2$
	SO_2NHCOPh	$\text{CH}_2\text{N(CH}_2\text{CH}_2)_2\text{O}$
	SO_2NHCOPh	OCH_2CF_3
	SO_2NHCOPh	$\text{OCH(CH}_3)_2$
	SO_2NHCOPh	SCH_2CH_3

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5. The compound of Claim 4 or a pharmaceutically acceptable salt thereof selected from the group consisting of:

5 5,7-dimethyl-2-ethyl-3[[2'-(N-n-pentylcarbonylsulfonamido)-5'-n-propyl[1,1'biphenyl]-4-yl]methylimidazo[4,5-b]pyridine;

10 5,7-dimethyl-2-ethyl-3[[2'-(N-n-butylaminocarbonylsulfonamido)-5'-n-propyl-[1,1'biphenyl]-4-yl]methylimidazo[4,5-b]pyridine;

15 5,7-dimethyl-2-ethyl-3[[2'-(N-(2-phenylethyl)carbonylsulfonamido)-5'-n-propyl[1,1'biphenyl]-4-yl]methylimidazo[4,5-b]pyridine;

20 5,7-dimethyl-2-ethyl-3[[2'-(N-(2-phenoxyphenyl)carbonylsulfonamido)-5'-n-propyl[1,1'biphenyl]-4-yl]methylimidazo[4,5-b]pyridine;

25 5,7-dimethyl-2-ethyl-3[[2'-(N-benzenecarbonylsulfonamido)-5'-n-propyl[1,1'biphenyl]-4-yl]methylimidazo[4,5-b]pyridine;

30 5,7-dimethyl-2-ethyl-3[[2'-(N-n-pentylcarbonylsulfonamido)-5'-dimethylaminomethyl [1,1'biphenyl]-4-yl]methylimidazo[4,5-b]pyridine;

5,7-dimethyl-2-ethyl-3[[2'-(N-n-butylaminocarbonylsulfonamido)-5'-n-butoxy[1,1'biphenyl]-4-yl]methylimidazo[4,5-b]pyridine;

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5,7-dimethyl-2-ethyl-3[[2'-(N-n-pentylcarbonylsulfon-
amido)-5'-isopropyl[1,1'biphenyl]-4-yl]methylimidazo
[4,5-b]pyridine;

5 5,7-dimethyl-2-ethyl-3[[2'-(N-ethoxymethylcarbonylsul-
fonamido)-5'-n-propyl[1,1'biphenyl]-4-yl]methylimidazo
[4,5-b]pyridine;

10 5,7-dimethyl-2-ethyl-3[[2'-(N-(n-butoxy)methylcarbonyl-
sulfonamido)-5'-n-propyl[1,1'biphenyl]-4-yl]methylini-
dazo[4,5-b]pyridine;

15 5,7-dimethyl-2-ethyl-3[[2'-(N-n-pentylcarbonylsulfon-
amido)-5'-pyrrolidin-1-ylmethyl[1,1'biphenyl]-4-yl]
methylimidazo[4,5-b]pyridine;

20 5,7-dimethyl-2-ethyl-3[[2'-(N-n-pentylcarbonylsulfon-
amido)-5'-morpholin-1-ylmethyl[1,1'biphenyl]-4-yl]
methylimidazo[4,5-b]pyridine; and

25 5,7-dimethyl-2-ethyl-3[[2'-(N-n-pentylcarbonylsulfon-
amido)-5'-n-butoxy[1,1'biphenyl]-4-yl]methylimidazo
[4,5-b]pyridine.

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6. A pharmaceutical composition useful in
the treatment of hypertension which comprises a
pharmaceutically acceptable carrier and a
pharmaceutically effective amount of a compound of
30 Claim 1 or a pharmaceutically acceptable salt thereof.

7. The composition of Claim 6 which
includes another antihypertensive selected from a

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diuretic selected from hydrochlorothiazide, chlorothiazide, chlorthalidone, methyclothiazide, furosemide, ethacrynic acid, triamterene, amiloride and spironolactone; a calcium channel blocker, selected from diltiazem, felodipine, nifedipine, nitrendipine and verapamil; a β -adrenergic antagonist selected from timolol, atenolol, metoprolol, propranolol, nadolol and pindolol; an angiotensin converting enzyme inhibitor selected from enalapril, lisinopril, captopril, ramipril, quinapril and zofenopril; a renin inhibitor selected from A-69729 and FK 744; an α -adrenergic antagonist selected from prazosin, doxazosin, and terazosin; a sympatholytic agent selected from methyldopa, clonidine and guanabenz; the atriopeptidase inhibitor, UK-79300; the serotonin antagonist, ketanserin; the A_2 -adenosine receptor agonist CGS 22492C; a potassium channel agonist selected from pinacidil and cromakalim; or another antihypertensive drug selected from reserpine, minoxidil, guanethidine, hydralazine hydrochloride and sodium nitroprusside; or combinations of the above-named drugs.

8. A method of treating hypertension which comprises administering to a patient in need of such treatment a pharmaceutically effective amount of a compound of Claim 1 or a pharmaceutically acceptable salt thereof.

9. An ophthalmological formulation for the treatment of ocular hypertension comprising an ophthalmologically acceptable carrier and an effective ocular antihypertensive amount of a

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compound of Claim 1 or a pharmaceutically acceptable salt thereof.

5 10. A method of treating ocular hypertension comprising administering to a patient in need of such treatment an effective ocular antihypertensive amount of a compound of Claim 1 or a pharmaceutically acceptable salt thereof.

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INTERNATIONAL SEARCH REPORT

International application No.

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A. CLASSIFICATION OF SUBJECT MATTER

IPC(5) :A61K 31/44; C07D 471/14, 413/10

US CL :546/118, 544/127, 514/303

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 546/118, 544/127, 514/303

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

CAS online, structure searched

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	EP, A, 0,399,731 (Roberts et al) 28 November 1990, see entire document.	1 to 10
Y	US, A, 5,128,327 (Chakravarty et al) 07 July 1992, see entire document.	1 to 10
Y	US, A, 5,102,880 (Chakravarty et al) 07 April 1992, see entire document.	1 to 10

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	*T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
*A document defining the general state of the art which is not considered to be part of particular relevance	*X documents of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
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*L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*g document member of the same patent family
*O document referring to an oral disclosure, use, exhibition or other means	
*P document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

21 October 1993

Date of mailing of the international search report

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Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231

Authorized officer

PHYLLIS SPIVACK

Facsimile No. NOT APPLICABLE

Telephone No. (703) 308-1235